WIT: integrated system for high-throughput genome sequence analysis and metabolic reconstruction

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ABSTRACT

The WIT (What Is There) (http://wit.mcs.ani.gov/WIT2/) system has been designed to support comparative analysis of sequenced genomes and to generate metabolic reconstructions based on chromosomal sequences and metabolic modules from the EMP/MPW family of databases. This system contains data derived from about 40 completed or nearly completed genomes. Sequence homologies, various ORFclustering algorithms, relative gene positions on the chromosome and placement of gene products in metabolic pathways (metabolic reconstruction) can be used for the assignment of gene functions and for development of overviews of genomes within WIT. The integration of a large number of phylogenetically diverse genomes in WIT facilitates the understanding of the physiology of different organisms.

INTRODUCTION

Starting with *Haemophilus influenza* (1) in 1995, over 20 microbial organisms have had their total genomic DNA sequenced and almost 100 others have been started as shown in the GOLD database (2). Currently we are observing an impressive development of the human genome project (3,4). In response to this growing amount of sequence data, computational tools for genome analysis have been developed and merged into shared analytical environments, such as GeneQuiz (5), KEGG (6), Pedant (7) and Entrez Genomes (8), moving cross-genome analysis to a new level. The development of analytical systems, together with the growth of sequencing data, have increased gene recognition rates from <50% (9,10) to >70% (11,12). Today, this remaining 30%, so-called 'hypothetical' or 'orphan' genes, separates us from a complete description of the genomic content and functions of an organism.

Computational approaches based on various types of clustering of potential genes, whether in phylogenetic space, as clusters of orthologous genes (COGs) (13) or position on the chromosome, such as in operons (14), increase the gene assignment level even further. An important stage of genome analysis is the integration of gene assignments into an organism-specific overview via so-called functional reconstruction (15), which is

the conceptual assembly of metabolic pathways, transport units and signal transduction pathways. It allows reconciliation of inconsistencies between different types of analysis, and often results in changes of initial gene function assignments based on similarity scoring.

The WIT system, discussed in this paper, represents the development of a genome analysis strategy in a multi-genome environment, which combines a variety of tools, dealing with individual open reading frames (ORFs) or proteins, with the ability to derive general conclusions. Using the WIT genome analysis system, a major part of the central metabolism of an organism can be reconstructed entirely *in silico* (16).

WIT: A VIEW TO A GENOME

The current version of the WIT system is available at Argonne National Laboratory (http://wit.mcs.anl.gov/WIT2) or at Integrated Genomics Inc. (http://wit.IntegratedGenomics.com/IGwit) and contains 43 complete or nearly complete genomes (Table 1).

These genomes consist of 123 482 predicted ORFs, of which 78 144 could be given functional assignments and 41 742 could be assembled into metabolic pathways, which came from EMP/MPW database (15). Pathways involved in the metabolism of carbohydrates and amino acids are connected into schematic overviews allowing the user to reveal substrates and final products connecting metabolic modules.

In order to incorporate a genome into WIT, a gene-searching program called CRITICA (17) can be used. Potential coding regions recognized in the DNA contigs are subjected to a FASTA search against the non-redundant database of assigned genes and loaded into the WIT system, together with the precomputed tables of best hits.

WIT provides a set of tools for the characterization of gene structures and functions, such as Functional Coupling, or Preserved Operons. WIT also provides integrated WWW access to such tools as PSI-BLAST, PROSITE, ProDom, COG, ClustalW and others. Functional content may be queried, for example, by looking for specific functions missing in the metabolic pathways, or by separating alternative gene functions derived from similarities found for a putative gene.

After genes have been assigned initial functions, they are then 'attached' to pathways by choosing templates from metabolic database (MPW) which best incorporate all observed functions. For any given organism, this usually leads to identification of

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Table 1. Genomes in WIT

Eukarya	Saccharomyces cerevisiae, Caenorhabditis elegans
Archaea	Sulfolobus solfataricus, Archaeoglobus fulgidus, Halobacterium sp., M.thermoautotrophicum, M.jannaschii, Pyrococcus furiosus, Pyrococcus horikoshii
Bacteria	A.aeolicus, C.trachomatis, Synechocystis sp., P.gingivalis, M.leprae, M.tuberculosis, B.subtilis, C.acetobutylicum, E.faecalis, M.genitalium, M.pneumoniae, S.pneumoniae, S.pyogenes, Rhizobium sp., R.capsulatus, S.aromaticivorans, N.gonorrhoeae, N.meningitidis, C.jejuni, H.pylori, E.coli, Y.pestis, H.influenzae, P.aeruginosa, B.burgdorferi, T.pallidum, D.radiodurans
Additional Genomes on the public server at Integrated Genomics Inc.	A.pernix, M.bovis, C.tepidum, S.typhi, T.maritima, A.actinomycetemcomitans, E.nidulans, Oryza sativa, A thaliana, R.prowazekii, P.abysii, C.pneumoniae, C.reinhardtii

functional sub-systems, as a model for further refinement. For example, it is now possible to identify inconsistencies, potentially missing enzymes/ORFs, thereby refining the model. When a basic model has been created, a curator finally evaluates this model against biochemical data and phenotypes known from the literature. The models come in both textual and graphical representations, fully linked with all underlying data. We call this whole process metabolic reconstruction, and the main role of the WIT system is to support this effort.

To examine or curate a functional model of an organism, one can use functions such as: Compare assignments, Summary of asserted functions and pathways, Examine trimmed ortholog clusters, Examine COG/trimmed ortholog cluster relationships, Search for pathways by regular expression, Search ORF functions by regular expression, Search ORF sequences by similarity search, Find NCBI's MEDLINE-references by EC-number, Search EMP by EC-number, and Find common proteins for organisms. Chromosomal clustering of functionally related genes (14) is another powerful component of the system, which recently allowed us to propose a number of candidate ORFs for 'orphan' metabolic functions. Continuous integration of newly sequenced genomes increases the depth of functional description by a reiterative process.

GAPPED GENOMES IN WIT

An important feature of the WIT system is its emphasis on incomplete or gapped genomes. Algorithms used for gene assignments depend on the size of a dataset used to cluster properties of ORFs, whether it is chromosomal position or ortholog clustering based on bi-directional best hits. By the incorporation of gapped genomes, even the public version of

WIT has integrated twice as much data as can be collected from only the completed genomes.

We believe that integrating systems like WIT can offer a solution for the problem of efficient use of incomplete sequence data. The gapped sequence contains a piece of almost every ORF, which allows the assignment of functions to almost all ORFs and the accurate reconstruction of the metabolism of the organism; good informatics can compensate for poorer sequence quality. A comparison of the results of analysis of the gapped genome of *Pseudomonas aeruginosa* with the complete genomes of *Escherichia coli* and *Bacillus subtilis* proves this statement (Table 2).

CONCLUSIONS

WIT has been designed to extract functional content from genome sequences and organize it into a coherent system, in order to facilitate post-sequencing experimental biology. The WIT system provides a set of local tools, which can be used to investigate functions of individual ORFs, based on similarities, motifs and various types of ORF clustering. It also generates overviews of functional subsystems and means to connect them into a complete picture of cellular functionality.

The WIT system is undergoing constant improvements, which can be traced in the PUMA-WIT-WIT2 line of development, and we believe that numerous further additions are needed to provide an adequate toolbox for the biological research community. Major directions of the ongoing WIT development are the following: (i) integration of structural data, which are currently underutilized in WIT; (ii) further development of the collection of functional maps and construction of more abstract scalable overviews, which should eventually cover all cellular functionality, and; (iii) development of a framework, which

Table 2. Comparison of the gapped P.aeruginosa genome with those of E.coli K-12 and B.subtilis 168

	Pseudomonas aeruginosa	Escherichia coli	Bacillus subtilis
Genome Size (Mb)	6.2	4.7	4.1
DNA assembled (%)	99	100	100
Total ORFs	5627	4289	4083
Assigned ORFs	4191	3499	3016
Asserted pathways	581	906	782
Missing assignments	133	102	178
No sequences	115	233	173

will integrate a flood of the differential display expression array data into the metabolic context.

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Improved tools for biological sequence comparison

(amino acid/nucleic acid/data base searches/local similarity)

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We have developed three computer pro-**ABSTRACT** grams for comparisons of protein and DNA sequences. They can be used to search sequence data bases, evaluate similarity scores, and identify periodic structures based on local sequence similarity. The FASTA program is a more sensitive derivative of the FASTP program, which can be used to search protein or DNA sequence data bases and can compare a protein sequence to a DNA sequence data base by translating the DNA data base as it is searched. FASTA includes an additional step in the calculation of the initial pairwise similarity score that allows multiple regions of similarity to be joined to increase the score of related sequences. The RDF2 program can be used to evaluate the significance of similarity scores using a shuffling method that preserves local sequence composition. The LFASTA program can display all the regions of local similarity between two sequences with scores greater than a threshold, using the same scoring parameters and a similar alignment algorithm; these local similarities can be displayed as a "graphic matrix" plot or as individual alignments. In addition, these programs have been generalized to allow comparison of DNA or protein sequences based on a variety of alternative scoring matrices.

We have been developing tools for the analysis of protein and DNA sequence similarity that achieve a balance of sensitivity and selectivity on the one hand and speed and memory requirements on the other. Three years ago, we described the FASTP program for searching amino acid sequence data bases (1), which uses a rapid technique for finding identities shared between two sequences and exploits the biological constraints on molecular evolution. FASTP has decreased the time required to search the National Biomedical Research Foundation (NBRF) protein sequence data base by more than two orders of magnitude and has been used by many investigators to find biologically significant similarities to newly sequenced proteins. There is a trade-off between sensitivity and selectivity in biological sequence comparison: methods that can detect more distantly related sequences (increased sensitivity) frequently increase the similarity scores of unrelated sequences (decreased selectivity). In this paper we describe a new version of FASTP, FASTA, which uses an improved algorithm that increases sensitivity with a small loss of selectivity and a negligible decrease in speed. We have also developed a related program, LFASTA, for local similarity analyses of DNA or amino acid sequences. These programs run on commonly available microcomputers as well as on larger machines.

METHODS

The search algorithm we have developed proceeds through four steps in determining a score for pair-wise similarity.

FASTP and FASTA achieve much of their speed and selectivity in the first step, by using a lookup table to locate all identities or groups of identities between two DNA or amino acid sequences during the first step of the comparison (2). The ktup parameter determines how many consecutive identities are required in a match. For example, if ktup = 4 for a DNA sequence comparison, only those identities that occur in a run of four consecutive matches are examined. In the first step, the 10 best diagonal regions are found using a simple formula based on the number of ktup matches and the distance between the matches without considering shorter runs of identities, conservative replacements, insertions, or deletions (1, 3).

In the second step of the comparison, we rescore these 10 regions using a scoring matrix that allows conservative replacements and runs of identities shorter than ktup to contribute to the similarity score. For protein sequences, this score is usually calculated using the PAM250 matrix (4), although scoring matrices based on the minimum number of base changes required for a replacement or on an alternative measure of similarity can also be used with FASTA. For each of these best diagonal regions, a subregion with maximal score is identified. We will refer to this region as the "initial region"; the best initial regions from Fig. 1A are shown in Fig. 1B.

The FASTP program uses the single best scoring initial region to characterize pair-wise similarity; the initial scores are used to rank the library sequences. FASTA goes one step further during a library search; it checks to see whether several initial regions may be joined together. Given the locations of the initial regions, their respective scores, and a "joining" penalty (analogous to a gap penalty), FASTA calculates an optimal alignment of initial regions as a combination of compatible regions with maximal score. FASTA uses the resulting score to rank the library sequences. We limit the degradation of selectivity by including in the optimization step only those initial regions whose scores are above a threshold. This process can be seen by comparing Fig. 1B with Fig. 1C. Fig. 1B shows the 10 highest scoring initial regions after rescoring with the PAM250 matrix; the best initial region reported by FASTP is marked with an asterisk. Fig. 1C shows an optimal subset of initial regions that can be joined to form a single alignment.

In the fourth step of the comparison, the highest scoring library sequences are aligned using a modification of the optimization method described by Needleman and Wunsch (5) and Smith and Waterman (6). This final comparison considers all possible alignments of the query and library sequence that fall within a band centered around the highest scoring initial region (Fig. 1D). With the FASTP program, optimization frequently improved the similarity scores of related sequences by factors of 2 or 3. Because FASTA calculates an initial similarity score based on an optimization of initial regions during the library search, the initial score is

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Abbreviation: NBRF, National Biomedical Research Foundation.

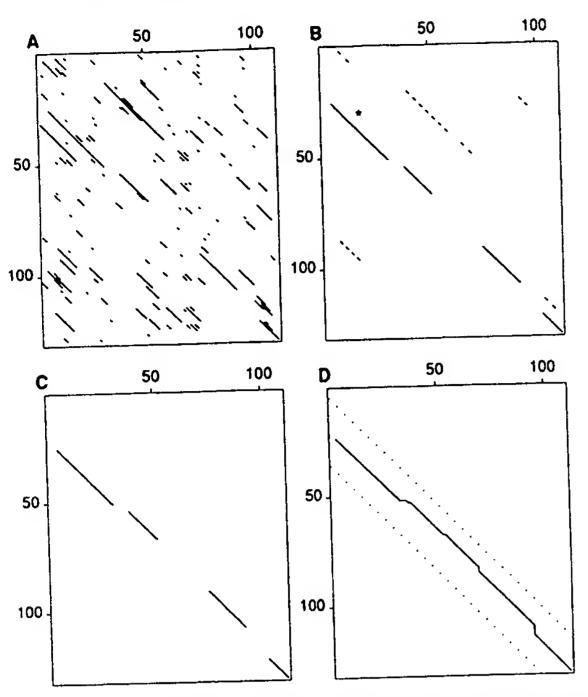


Fig. 1. Identification of sequence similarities by FASTA. The four steps used by the FASTA program to calculate the initial and optimal similarity scores between two sequences are shown. (A) Identify regions of identity. (B) Scan the regions using a scoring matrix and save the best initial regions. Initial regions with scores less than the joining threshold (27) are dashed. The asterisk denotes the highest scoring region reported by FASTP. (C) Optimally join initial regions with scores greater than a threshold. The solid lines denote regions that are joined to make up the optimized initial score. (D) Recalculate an optimized alignment centered around the highest scoring initial region. The dotted lines denote the bounds of the optimized alignment. The result of this alignment is reported as the optimized score.

much closer to the optimized score for many sequences. In fact, unlike FASTP, the FASTA method may yield initial scores that are higher than the corresponding optimized scores.

Local Similarity Analyses. Molecular biologists are often interested in the detection of similar subsequences within longer sequences. In contrast to FASTP and FASTA, which report only the one highest scoring alignment between two sequences, local sequence comparison tools can identify multiple alignments between smaller portions of two sequences. Local similarity searches can clearly show the results of gene duplications (see Fig. 2) or repeated structural features (see Fig. 3) and are frequently displayed using a "graphic matrix" plot (7), which allows one to detect regions of local similarity by eye. Optimal algorithms for sensitive local sequence comparison (6, 8, 9) can have tremendous computational requirements in time and memory, which make them impractical on microcomputers and, when comparing longer sequences, on larger machines as well.

The program for detecting local similarities, LFASTA, uses the same first two steps for finding initial regions that FASTA uses. However, instead of saving 10 initial regions, LFASTA saves all diagonal regions with similarity scores greater than a threshold. LFASTA and FASTA also differ in the construction of optimized alignments. Instead of focusing on a single region, LFASTA computes a local alignment for each initial region. Thus LFASTA considers all of the initial regions shown in Fig. 1B, instead of just the diagonal shown in Fig. 1D. Furthermore, LFASTA considers not

only the band around each initial region but also potential sequence alignments for some distance before and after the initial region. Starting at the end of the initial region, an optimization (6) proceeds in the reverse direction until all possible alignment scores have gone to zero. The location of the maximal local similarity score in the reverse direction is then used to start a second optimization that proceeds in the forward direction. An optimal path starting from the forward maximum is then displayed (5). The local homologies can be displayed as sequence alignments (see Fig. 2B) or on a two-dimensional graphic matrix style plot (see Figs. 2A and 3).

Statistical Significance. The rapid sequence comparison algorithms we have developed also provide additional tools for evaluating the statistical significance of an alignment. There are approximately 5000 protein sequences, with 1.1 million amino acid residues, in the NBRF protein sequence library, and any computer program that searches the library by calculating a similarity score for each sequence in the library will find a highest scoring sequence, regardless of whether the alignment between the query and library sequence is biologically meaningful or not. Accompanying the previous version of FASTP was a program for the evaluation of statistical significance, RDF, which compares one sequence with randomly permuted versions of the potentially related sequence.

We have written a new version of RDF (RDF2) that has several improvements. (i) RDF2 calculates three scores for each shuffled sequence: one from the best single initial region (as found by FASTP), a second from the joined initial regions (used by FASTA), and a third from the optimized diagonal. (ii) RDF2 can be used to evaluate amino acid or DNA sequences and allows the user to specify the scoring matrix to be employed. Thus sequences found using the PAM250 scoring matrix can be evaluated using the identity or genetic code matrix. (iii) The user may specify either a global or local shuffle routine.

Locally biased amino acid or nucleotide composition is perhaps the most common reason for high similarity scores of dubious biological significance (10). High scoring alignments between query and library sequences may be due to patches of hydrophobic or charged amino acid residues or to A+T- or G+C-rich regions in DNA. A simple Monte Carlo shuffle analysis that constructs random sequences by taking each residue in one sequence and placing it randomly along the length of the new sequence will break up these patches of biased composition. As a result, the scores of the shuffled sequences may be much lower than those of the unshuffled sequence, and the sequences will appear to be related. Alternatively, shuffled sequences can be constructed by permuting small blocks of 10 or 20 residues so that, while the order of the sequence is destroyed, the local composition is not. By shuffling the residues within short blocks along the sequence, patches of G+C- or A+T-rich regions in DNA, for example, are undisturbed. Evaluating significance with a local shuffle is more stringent than the global approach, and there may be some circumstances in which both should be used in conjunction. Whereas two proteins that share a common evolutionary ancestor may have clearly significant similarity scores using either shuffling strategy, proteins related because of secondary structure or hydropathic profile may have similarity scores whose significance decreases dramatically when the results of global and local shuffling are compared.

Implementation. The FASTA/LFASTA package of sequence analysis tools is written in the C programming language and has been implemented under the Unix, VAX/VMS, and IBM PC DOS operating systems. Versions of the program that run on the IBM PC are limited to query se-

Table 1. FASTA and FASTP initial scores of the T-cell receptor (RWMSAV) versus the NBRF data base

	Init		score
NBRF code	Sequence	FASTA	FASTP
RWHUAV	T-cell receptor α chain	155	98
K1HURE	Ig k chain V-I region	127	111
KVMS50	Ig k chain V region	149	62
KVMSM6	Ig k chain precursor V regions	141	64
KVRB29	Ig k chain V region	126	54
L3HUSH	Ig λ chain V-III region	90	47
KVMS41	Ig k chain precursor V region	87	87
RWMSBV	T-cell receptor β -chain precursor	94	94
RWHUVY	T-cell receptor β -chain precursor	91	59
RWHUGV	T-cell receptor γ -chain precursor	87	61
RWHUT4	T-cell surface glycoprotein T4	86	63
RWMSVB	T-cell receptor γ -chain precursor	71	41
HVMS44	Ig heavy-chain V region	67	36
G1HUDW	Ig heavy-chain V-II region	62	35

The average FASTP score = 26.1 ± 6.8 (mean \pm SD). The average FASTA score = 26.2 ± 7.2 (mean \pm SD). The mean and SD were computed excluding scores >54. V, Variable.

quences of 2000 residues; library sequences can be any length. Copies of the program are available from the authors.

Although FASTA and LFASTA were designed for protein and DNA sequence comparison, they use a general method that can be applied to any alphabet with arbitrary match/mismatch scoring values. All the scoring parameters, including match/mismatch values, values for the first residue in a gap and subsequent residues in the gap, and other parameters that control the number of sequences to be saved and the histogram intervals, can be specified without changing the program.

EXAMPLES

Comparison of FASTA with FASTP. To demonstrate the superiority of the FASTA method for computing the initial score, we compared the protein sequence of a T-cell receptor α chain (NBRF code RWMSAV) with all sequences in the NBRF protein data base[‡] and computed initial scores with both the present and previous methods. The T-cell receptor is a member of the immunoglobulin superfamily; in Release 12.0 of the data base, this superfamily has 203 members. FASTP placed 160 immunoglobulin superfamily sequences in the 200 top-scoring sequences; 57 related sequences received initial scores less than four standard deviations above the mean score. FASTA placed 180 superfamily members in the 200 top-scoring sequences; only 20 related sequences scored below four standard deviations above the mean. Table 1 contains specific examples from this data base search. Although there is often little difference in the two methods, this example shows that in a number of cases the new method obtains significantly higher scores between related sequences.

Nucleic Acid Data Base Search. FASTA can also be used to search DNA sequence data bases, either by comparing a DNA query sequence to the DNA library or by comparing an amino acid query sequence to the DNA library by translating each library DNA sequence in all six possible reading frames. We compared the 660-nucleotide rat transforming growth factor type α mRNA (GenBank locus RATTGFA) with all the mammalian sequences in Release 48 of GenBank§. We set ktup = 4 (see Methods), and the search was completed in under 15 min on an IBM PCAT microcom-

Table 2. DNA data base search of rat transforming growth factor (RATTGFA) versus mammalian sequences

GenBank	GenBank		Score
locus	Sequence	Initial	Optimized
HUMTFGAM	Human TGF mRNA	1336	1618
HUMTGFA2	Human TGF gene (exon 2)	354	366
HUMTGFA1	Human TGF gene (5' end)	224	381
MUSRGEB3	Mouse 18S-5.8S-28S rRNA gene	140	107
MUSRGE52	Mouse 18S-5.8S-28S rRNA gene	140	107
MUSMHDD	MHC class I H-2D	122	78
HUMMETIF1	Metallothionein (MT)I _F gene	116	92
MUSRGLP	45S rRNA (5' end)	115	83
HUMPS2	pS2 mRNA	105	106
MUSC1AI1	α-1 type I procollagen	86	89

The 10 sequences having the highest initial scores are given. TGF, transforming growth factor; MHC, major histocompatibility complex.

puter. The 10 top-scoring library sequences are shown in Table 2. Although it can be seen that the 3 top-scoring sequences are clearly related to RATTGFA, there are other high-scoring sequences that are probably not related, and the mouse epidermal growth factor, found in the translated data base search (Table 3), is not found among the top-scoring sequences.

To further examine the similarity detected between RAT-TGFA and MUSRGEB3, a mouse rRNA gene cluster, we used the RDF2 program for Monte Carlo analysis of statistical significance (the window for local shuffling was set to 10 bases). Of the 50 shuffled comparisons (data not shown), 1 obtained an initial score greater than 140 (the observed initial score), and 9 shuffled sequences obtained optimized scores greater than 107 (the observed optimized score). Therefore, the similarity between RATTGFA and MUSRGEB3 is unlikely to be significant.

Translated Nucleic Acid Data Base Search. When searching for sequences that encode proteins, amino acid sequence comparisons are substantially more sensitive than DNA sequence comparisons because one can use scoring matrices like the PAM250 matrix that discriminate between conservative and nonconservative substitutions. A variant of FASTA, TFASTA, can be used to compare a protein sequence to a DNA sequence library; it translates the DNA sequences into each of six possible reading frames "on-the-fly." TFASTA translates the DNA sequences from beginning to end; it includes both intron and exon sequences in the translated protein sequence; termination codons are translated into unknown (X) amino acids. Table 3 shows the results of a translating search of the mammalian sequences in the Gen-Bank DNA data base using the RATTGFA protein sequence as the query and ktup = 1. In the translated search, the mouse epidermal growth factor now obtains an initial score higher than any unrelated sequences; however, HUMTGFA1, which was found in the DNA data base search but only contains 13 translated codons, is no longer among the top scoring sequences.

Local Similarities. Fig. 2 displays the output of a local similarity analysis (ktup = 4) of CHPHBA1M, a chimpanzee α 1-globin mRNA, and RABHBAPT, a rabbit α -globin gene, including the complete coding sequence and a flanking pseudo- θ_1 -globin gene. LFASTA can either display a graphic matrix style plot of the local homologies (Fig. 2A) or the alignments themselves (Fig. 2B). The right-most three alignments (Fig. 2A) match the corresponding regions of the mRNA to exon subsequences from the pseudogene. We note that the FASTA initial score for the comparison of CHPH-

Protein Identification Resource (1987) Protein Sequence Database (Natl. Biomed. Res. Found., Washington, DC), Release 12. §EMBL/GenBank Genetic Sequence Database (1987) (Intelligenetics, Mountain View, CA), Tape Release 48.

Table 3. Translated DNA data base search of rat transforming growth factor (RATTGFA) versus mammalian sequences

CanDonk		· · · · · · · · · · · · · · · · · · ·		Score
GenBank locus	Sequence	Frame	Initial	Optimized
RATTGFA	Rat TGF type α	1	816	816
HUMTGFAM	Human TGF mRNA	2	671	770
HUMTGFA2	Human TGF gene	1	204	205
MUSEGF	Mouse EGF mRNA	3	93	129
MUSMHAB3	Mouse MHC class II H2-IA _B	1	91	58
MUSIGCD17	Mouse Ig germ-line DJC region	3′	85	48
HUMESTR	Human estrogen receptor	3	83	65
RATINSI	Rat insulin 1 (Ins-1) gene	2	81	63
MUSTHYS1	Mouse thymidylate synthase	2	80	63
HUMPNU3	Human purine nucleoside phosphorylase	1'	80	52

The 10 sequences having the highest initial scores are given. TGF, transforming growth factor; EGF, epidermal growth factor; D, diversity; J, joining; C, constant; MHC, major histocompatibility complex.

BA1M and RABHBAPT would be based on the three globin gene exons, while the FASTP initial score would be based on a single conserved exon.

The Smith-Waterman optimization used in the LFASTA program allows the detection of more subtle features than can be detected by the eye using a graphic matrix plot, because the path traced is locally optimal, even though it may only have a slightly higher density of identities and conservative replacements. Fig. 3 shows a plot from a local similarity self-comparison of the myosin heavy chain from the nematode Caenorhabditis elegans (MWKW) using the PAM250 matrix. The amino-terminal half of the molecule forms a large globular head without any periodic structure; the solid line down the main diagonal represents the expected identity of the sequence with itself. The symmetrical parallel lines along the carboxyl-terminal half of the molecule correspond to the 28-residue repeat responsible for the α -helical coiled-coil structure of the rod segment.

DISCUSSION

In searching a data base, one is attempting to measure relatedness; in aligning two homologous sequences, one is

trying to choose the most likely set of mutations since their divergence from a common ancestral sequence. Thus any tool for the analysis of sequence similarities must contain within it an implicit model of molecular evolution. An algorithm that guarantees the optimality of its alignments based on a set of scoring rules must be judged on how well these rules fit our current understanding of the process of molecular evolution. Algorithms that sacrifice realism to achieve greater efficiency, regardless of their mathematical rigor, require careful empirical evaluation.

Even though the tools we have developed use rigorous algorithms at each step and incorporate a realistic model of evolution, their hierarchical nature make them heuristic. The original FASTP program has had the benefit of extensive use and evaluation by a wide variety of scientists. The FASTA program exploits refinements of the previous approach that result in a significant improvement in sensitivity. The LFA-STA local similarity analysis program is also a logical extension of the FASTP approach.

Because of the trade-offs between sensitivity and selectivity in data base searches, the results of any search, and particularly those that result in alignment scores that are not clearly separated from the distribution of all library sequence

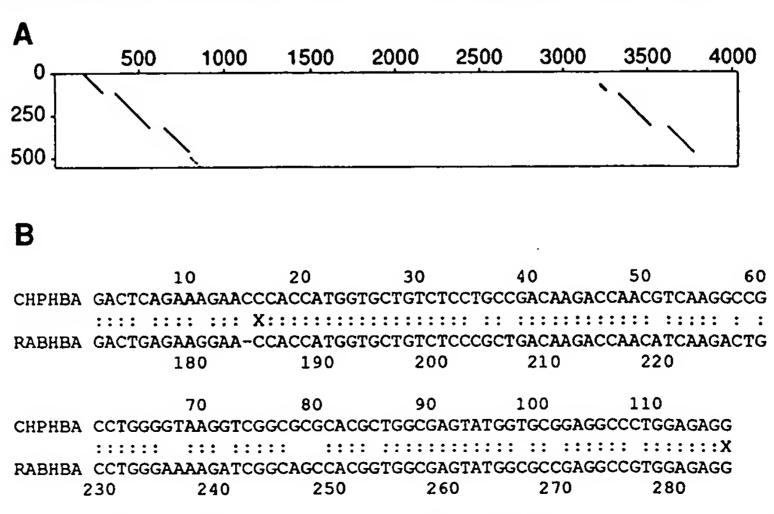
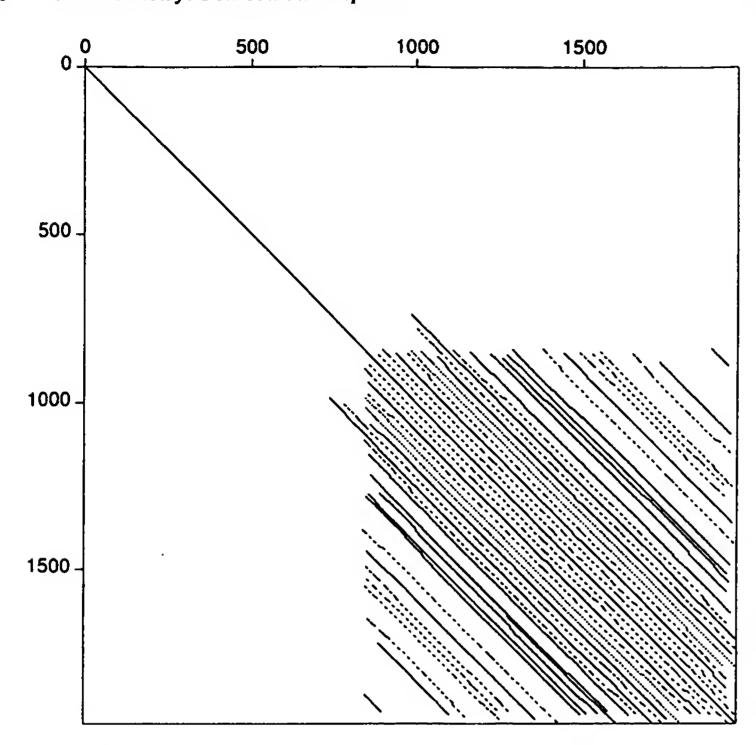


Fig. 2. Local comparison of an α -globin mRNA sequence with an α -globin gene cluster. An ape α_1 -globin mRNA sequence (GenBank sequence CHPHBA1M) was compared with a rabbit α -globin gene sequence (RABHBAPT) containing a second pseudo- θ -globin gene using the LFASTA program. (A) A plot of the homologous regions shared by the two sequences. (B) One of the alignments between the mRNA sequence and the rabbit α -globin gene (nucleotides 171-855). Three other alignments between the mRNA sequence and the α -globin gene and three alignments between the pseudo- θ -globin gene (nucleotides 3200-3770) were calculated but are not shown. There is 84.3% identity in the 115 nucleotide overlap. The initial region and optimized scores using LFASTA are 284 and 304, respectively. X denotes the ends of the initial region found by LFASTA.



Repeated structure in the myosin heavy chain. LFASTA was used to compare the Caenorhabditis elegans myosin heavy chain protein sequence (NBRF code MWKW) with itself using the PAM250 scoring matrix. The solid, dashed, and dotted lines denote decreasing similarity scores. The solid lines had initial region scores greater than 80 and optimized local scores greater than 150; the longer dashed lines had initial region and optimized local scores greater than 65 and 120, respectively, and the shorter dashed lines had initial region and optimized local scores greater than 50 and 100, respectively. Homologous regions with lower scores are plotted with dots.

scores, must be carefully evaluated (1, 11). The Monte Carlo analysis of statistical significance provided by a program such as RDF2 can often be critical in evaluating a borderline similarity. Previously we suggested ranges of z values [(observed score - mean of shuffled scores)/standard deviation of shuffled scores] corresponding to approximate significance levels. However the z values determined in a Monte Carlo analysis become less useful as the distribution of shuffled scores diverges from a normal distribution, as is found with FASTA. Therefore, we now focus on the highest scores of the shuffled sequences. For example, if in 50 shuffled comparisons, several random scores are as high or higher than the observed score, then the observed similarity is not a particularly unlikely event. One can have more confidence if in 200 shuffled comparisons, no random score approaches the observed score. In general, our experience has led us to be conservative in evaluating an observed similarity in an unlikely biological context.

These programs provide a group of sequence analysis tools that use a consistent measure for scoring similarity and constructing alignments. FASTA, RDF2, and LFASTA all use the same scoring matrices and similar alignment algorithms, so that potentially related library sequences discov-

ered after the search of a sequence data base can be evaluated further from a variety of perspectives. In addition, LFASTA can also show alternative alignments between sequences with periodic structures or duplications.

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APPENDIX C

Beyond complete genomes: from sequence to structure and function

Eugene V Koonin*, Roman L Tatusov and Michael Y Galperin

Computer analysis of complete prokaryotic genomes shows that microbial proteins are in general highly conserved — —70% of them contain ancient conserved regions. This allows us to delineate families of orthologs across a wide phylogenetic range and, in many cases, predict protein functions with considerable precision. Sequence database searches using newly developed, sensitive algorithms result in the unification of such orthologous families into larger superfamilies sharing common sequence motifs. For many of these superfamilies, prediction of the structural fold and specific amino acid residues involved in enzymatic catalysis is possible. Taken together, sequence and structure comparisons provide a powerful methodology that can successfully complement traditional experimental approaches.

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Abbreviations

COGs clusters of orthologous groups
HAD haloacid dehalogenase

Introduction

The determination of the complete genome sequences of several bacteria and archea and one eukaryote [1-6,7"-12"] marked the beginning of a new age in biology. For the first time, we can take a look at the complete set of proteins present in the cells of each particular organism and try to identify the proteins responsible for each cellular function. In cases where no known proteins can be found to perform a particular task, the most likely substitutes can be predicted from the set of unassigned gene products. Clearly this can be done only by analysis of complete genomes, as partial sequences do not allow us to ascertain that certain proteins are not encoded in a given genome [13]. These new approaches are gradually changing our understanding of a variety of biological phenomena. As the number of sequenced genomes is expected to grow exponentially for the next few years, their impact on different biological disciplines will increase. We have recently discussed the implications of the complete genomes for microbial evolution [14]. Here we consider the effect of the genome revolution, together with the improving methods for sequence analysis, on our ability to predict and understand protein structure and function.

Towards a natural taxonomy of proteins and protein families

The numerous genome sequencing projects have resulted in a rapid growth of protein databases (see, e.g. [15]). In contrast to the pre-genome era, when researchers typically chose to clone and sequence genes with documented functional roles, we are now getting many protein sequences whose functions are not known. This presents a challenge to extract the most from these sequences in terms of salient features of the encoded proteins, for example to classify them according to their homologous relationships, and to predict their possible catalytic activities and/or cellular functions, three-dimensional (3D) structures and evolutionary origin.

Protein classifications, pioneered by Dayhoff and her coworkers, have historically been based on sequence alignments. Similar proteins formed families, which were combined into superfamilies [16]. This approach, continued in the PIR database [17], proved extremely popular. However, even PIR superfamilies often unite closely related proteins and more distant relationships are being missed. Other protein databases, such as PROSITE [18], PRINTS [19], Pfam [20], and ProDom [21], group proteins on the basis of conserved sequence motifs and, generally, contain much more diverse protein families. Structural comparisons of proteins, implemented in FSSP, CATH and SCOP databases, offer yet another approach to protein classification [22-24]. SCOP superfamilies, for example, unite proteins that have some similarities in their 3D structures, but often no detectable sequence similarity [25]. Thus, in the absence of clear sequence or structural similarities, the criteria for inclusion of distantly related proteins into a family (or superfamily) become increasingly arbitrary.

With the inception of extensive genome sequencing, it has become possible to classify genes and proteins on a different principle, namely by delineating families of paralogs—related genes within the same genome [26,27]. Such analyses have revealed a complex hierarchical organization of paralogous families in each of the studied genomes and produced at least two generalizations: first, the fraction of genes that belong to families of paralogs increases with the increase of the total number of genes in a genome: from ~25% in the minimal genome of Mycoplasma genitalium to >50% in the large (for a prokaryote) Escherichia coli genome; second, the largest superfamilies of paralogs are mostly the same in all genomes [28–33].

Knowledge of all the protein sequences from multiple complete genomes (Table 1) allows us to redefine the entire Methanobacterium thermoautotrophicum

Archaeoglobus fulgidus

Caenorhabditis elegans

Saccharomyces cerevisiae

Table 1

Species	Proteins end	coded in the genome*	COGs found (% total)	3D s	Iructures
	Total number	Belong to COGs [†] (% total)		In PD8	Predicted*
Escherichia coli	4289	2003 (47%)	821 (95%)	240	887
Haemophilus influenzae	1717	979 (57%)	658 (77%)	2	267
Helicobacter pylori	1566	841 (54%)	617 (72%)	0	169
Synechocystis sp.	3169	1551 (49%)	796 (93%)	2	431
Borrelia burgdorferi	850	483 (57%)	363 (42%)	0	105
Bacillus subtilis	4100	· 1945 (47%)	732 (85%)	12	578
Mycoplasma genitalium	467	341 (75%)	290 (34%)	0	75/103
Mycoplasma pneumoniae	677	378 (56%)	309 (36%)	0	78
Methanococcus jannaschii	1715	830 (48%)	498 (58%)	0	170

1869

2407

5932

12,178

897 (48%)

1131 (47%)

1736 (29%)

2172 (18%)

problem of protein classification. Since the fraction of proteins conserved over large phylogenetic distances (ancient conserved domains) appears to be nearly constant at ~70% in all prokaryotic genomes [34], it becomes feasible to replace more or less arbitrary clustering of proteins by similarity with consistent groups in which the evolutionary relationships between the members are specifically defined. Such a classification of proteins can provide a framework for evolutionary studies and for rapid, largely automatic, functional annotation of newly sequenced genomes.

Several classifications of homologous proteins encoded in complete genomes have been produced, based on allagainst-all protein sequence comparisons [35,36,37"]. Each of these projects is aimed at the identification of orthologs. that is direct counterparts in different genomes, connected by an uninterrupted line of vertical descent and typically retaining their physiological function [26,27]. In particular, the system of clusters of orthologous groups (COGs) was designed to accommodate the vastly different evolution rates observed for different genes [37"]. The COGs construction procedure identifies the closest homologs in each of the sequenced genomes for each protein, even if the similarity is fairly low and not statistically significant by itself. The approach to the identification of COGs was built upon the transitivity of orthologous relationships, that is the simple notion that any group of at least three genes from distant genomes, which are more similar to each other than they are to any other genes from the same genomes, is most likely to belong to an orthologous family. Clearly, this is a probabilistic assumption based on a 'weak molecular clock concept', which posits that orthologs are more similar to each other than they are to paralogs with different, even if related, functions. This assumption, however, seems to hold true in cases where we have reasons to accept orthology on functional grounds (for example, aminoacyl-tRNA synthetases or ribosomal proteins). Orthology is not necessarily a one-to-one relationship, as in cases of lineage-specific duplications, orthology can only be established between families of paralogous genes. Such complex relationships require caution in the functional interpretation of the phylogenetic classification of proteins. Nevertheless, about 60% of the original set of 720 COGs [37"] are simple families, with no paralogs or with paralogs from one lineage only, suggesting the possibility of straightforward transfer of functional information from functionally characterized genes from model systems such as *E. coli* and yeast to those from poorly characterized genomes.

484 (56%)

512 (60%)

577 (67%)

466 (54%)

199

290

846

NA

45

The utility of this system of protein classification was tested on several newly sequenced bacterial, archeal and eukaryotic genomes. Interestingly, with the only exception of the minimal genome of M. genitalium, the fraction of the proteins that belong to the COGs — ancient families conserved across a wide phylogenetic range — is about the same and very close to 50% for all prokaryotic genomes (Table 1). This is clearly compatible with the previous estimate that about 70% of the proteins encoded in each genome contain ancient conserved regions. The fraction of the proteins included in the COGs is at this time lower, which is evidently due to the requirement for three distant lineages to be included, and to the limited number of species in the first instalment of the COGs. There is little doubt that with new genomes added, the number of COGs will asymptotically approach the total number of ancient conserved regions. By contrast, this fraction is much lower

^{*}The numbers are from the latest updates in the GenBank genome division (ftp://ncbi.nlm.nih.gov/genbank/genomes). C. elegans genome is about 85% complete; the data are from Wormpep12 (www.sanger.ac.uk/Projects/C_elegans/wormpep). †Based on the set of 860 COGs, obtained by adding H. pylori proteins to the original set of 720 COGs [37**]. †The numbers are from the PEDANT database [53*], calculated by comparing the protein set encoded in each genome to the PDB using FASTA with cutoff score of 120; the second figure for M. genitalium is from [54*]; the data for C. elegans are not available.

for eukaryotic genomes, indicating the prevalence of enkaryote-specific families.

Comparison of the new protein sets with the COGs resulted in a number of functional predictions for previously uncharacterized proteins. Even for the Helicobacter pylori proteins, most of which show highly significant similarity to homologs from E. coli and other bacteria and have been described in considerable detail [8"], predictions were made in more than 100 cases (http://www.ncbi.nlm.nih/COG); function was also predicted for a number of archeal and worm proteins (EV Koonin, RL Tatusov, MY Galperin, unpublished data).

Missing gene families and evolution of metabolic pathways

Comparative analysis of the available complete genomes shows that metabolic diversity generally correlates with genome size. Parasitic bacteria import a variety of metabolites, which allows them to shed genes encoding enzymes for many or even most of the metabolic pathways [1-3, 84,33,38]. In contrast, all cells have to rely on their own gene products for performing such essential functions as genome expression, replication and repair, and membrane biogenesis and others. These tasks alone require at least about 200 genes [13,37"].

Given complete genome sequences, classification of proteins into orthologous groups provides a convenient way to systematically survey the protein families present or absent in a genome and to identify the metabolic pathways that are likely to be operative in the organism analyzed. When some of the required enzymes cannot be found in the genome, the respective parhways are either not operative, or use other, unrelated, proteins to catalyze the missing steps (see [39]). An example of such an analysis, which included superposition of the phylogenetic patterns derived from the COGs [37"], over the scheme of glycolysis, reveals several interesting trends (Figure 1), Glycolysis includes three reactions that in different species are catalyzed by non-orthologous enzymes, namely phosphofructokinases, aldolases and phosphoglycerate mutases. Interestingly, the second phosphofructokinase in E. roli, encoded by the pfkB gene, has apparently been recruited from a ubiquitous family of ribokinase-like sugar kinases. The ribokinase COG seems to be an example of a complex family in which the exact orthologous connections are not always easy to trace. In particular, even though PfkB formally belongs to the COG, there seems to be no actual ortholog of it in other genomes. Thus H. pylori does not encode a phosphofructokinase at all, although it has genes for other kinases of the ribokinase family and, accordingly, is represented in the respective COG (Figure 1).

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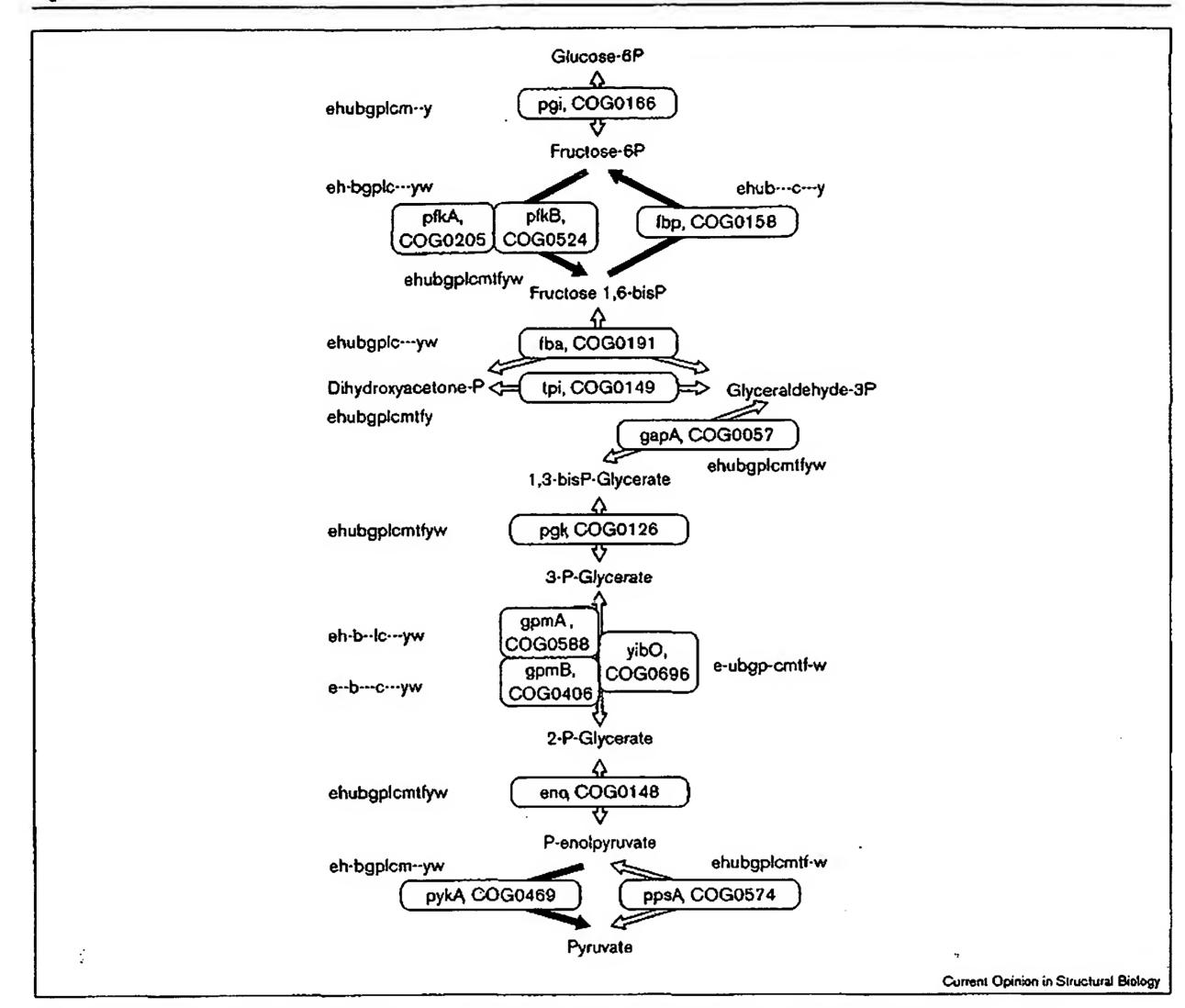
A remarkable case of non-orthologous gene displacement involves two unrelated forms of phosphoglycerate mutase, the 2,3-bisphosphoglycerate (BPG)-dependent and the BPG-independent one. While H. influenzae and Borrelia burgdorferi encode only the BPG-dependent form, and H. pylori, mycoplasmas, and archea encode only the BPGindependent form (see [40]), free-living bacteria such as E. coli, Bacillus subtilis and Synechocystis sp. possess genes coding for both these forms, with two paralogs of the BPGdependent one (Figure 1). Phosphofructokinase, aldolase and fructose bisphosphatase genes are all missing in the archea (Figure 1), in accordance with the experimental data [41]. This is consistent with the idea that glycolysis originally evolved as a biosynthetic pathway, containing only the lower (tri-carbon) part [42].

Systematic identification of missing links in functional systems in organisms for which complete genome sequences are available is probably the most important application of protein family classification. Conspicuous gaps in the H. pylori metabolism became apparent from the COG analysis, suggesting major revisions to the general scheme of the central metabolic pathways in this bacterium (Table 2). In particular, unlike most other bacteria (and all with completely sequenced genomes), H. pylori seems to possess neither glycolysis nor the pentose phosphate shunt, the Entner-Doudoroff pathway being the only major route of sugar catabolism. Indeed, sugar fermentation, resulting in intracellular acid production, would be an additional burden on the pH maintenance mechanism in this bacterium, which has to survive in an external pH of 2-3. By contrast, gluconeogenesis, which converts organic acids into sugars required for nucleic acid and peptidoglycan biosynthesis and thus removes H+ from the cytoplasm, appears to be fully functional in H. pylori. For the purpose of energy production, H. pylori apparently depends on amino acid fermentation, which causes alkalinization of the cytoplasm and thus relieves part of the problem of pH maintenance. Amino acids and oligopeptides that serve as substrates for this fermentation are produced by gastric proteolysis and transported by readily identifiable permeases.

From genomes and families to superfamilies and folds

Classification systems aimed at the identification of families of orthologs make no attempt to capture the more subtle conserved motifs in proteins, which reflect ancient relationships at the level of superfamilies and frequently are critically important for understanding protein functions and structures [43,44]. Computer methods for the detection of such motifs and delineation of superfamilies have lately progressed significantly through programs such as BLIMPS/MULTIMAT [45], Probe [46], and PSI-BLAST [47"], which combine pairwise sequence comparisons with profile analysis. PSI-BLAST, in particular, has proved to be a powerful tool for the detection of subtle sequence motifs, resulting in the discovery of a number of unsuspected superfamily relationships [47",48"]. Furthermore, one of the perhaps under-appreciated benefits of the accumulation of genomic sequences is the greatly improved capacity to identify even very subtle sequence similarities due to

Figure 1



Glycolytic enzymes in organisms with completely sequenced genomes. The enzymes are listed under E. coli gene names. The COG numbers are as in COG database (www.ncbi.nlm.nih.gov/COG, [37**]) (where available). Shaded arrows indicate reversible reactions, black arrows practically irreversible ones. Phosphoenolpyruvate synthase-catalyzed reaction in the direction of phosphoenolpyruvate hydrolysis has been demonstrated in vitro. Phylogenetic patterns are: e, Escherichia coli; h, Haemophilus influenzae; u, Helicobacter pylori; b, Bacillus subtilis; g, Mycoplasma genitalium; p, Mycoplasma pneumoniae; l, Borrelia burgdorferi; c, Synechocystis sp.; m, Methanococcus jannaschii; t, Methanobacterium thermoautotrophicum; f, Archaeoglobus fulgidus; y, Saccharomyces cerevisiae; w, Caenorhabditis elegans.

the increasingly uniform population of the protein universe by these relatively unbiased sequence sets, of which the new methods for sequence analysis mentioned above can take advantage [49].

In the past year, we have seen the identification or significant extension of a number of protein superfamilies; some examples, with the distribution among complete genomes, are shown in Table 3. Most of these superfamilies are universally found in all genomes, with the counts more or less proportional to the total number of genes in the genome. Some expansions are, however, remarkable,

such as, for example, urease-related hydrolases and ATP-grasp domains in the archea, and HAD superfamily hydrolases in *E. coli* and *B. subtilis* (Table 3). In certain cases, the phylogenetic distribution of a superfamily immediately suggests major evolutionary events. Thus the BRCT domain is present in a single copy in the DNA ligase of all bacteria (with one additional copy found only in *Synechocystis*), is missing in the archea, and is dramatically expanded in its distribution in the eukaryotes (Table 3). The most obvious interpretation of this distribution is that this domain has entered the eukaryotic world by horizontal gene transfer from bacteria and has undergone exten-

Table 2

Enzyme activity	E. coli gene	COG number	Status in H. pylori	Implications for H. pylori metabolism
Phosphofructokinase	plkA plkB	COG0206 COG0525	Missing Present (ribokinase)	Absence of the two key glycolytic enzymes shows that Embden-Meyerhof pathway is not functional in H. pylori.
Pyruvate kinase	pykA pykF	COG0470	Missing	Gluconeogenesis enzymes, bypassing these reactions, fructose bisphosphatase (HP1385) and phosphoenolpyruvate synthase (HP0121), are present in H. pylori, allowing it to produce sugars required for peptidoglycan biosynthesis.
6-phosphogluconate dehydrogenase	gnd	COG0360	Missing	Pentose phosphate pathway is also not functional. Even though <i>H. pylori</i> has a ribose 5-phosphate isomerase
Ribose 5-phosphate isomerase	rpiA	COG0120	Missing	encoded by an ortholog of the E. coli <i>rpiB</i> , no gene coding for 6-phosphogluconate dehydrogenase could be identified. The only saccharolytic pathway in <i>H. pylori</i> appears to be the Entner-Doudoroff pathway.
Lipoate synthase	liρA	COG0318	Missing	Pyruvate dehydrogenase complex is absent in H. pylori;
Lipoate-protein	<i>lpIA</i>	COG0411	Missing	acetate kinase and phosphotransacetylase are not
ligase	lipB	COG0319	Missing	functional. Pyruvate-ferredoxin oxidoreductase is the only
Dihydrolipoamide acyltransferase	aceF	COG0510	Missing	acetyl-CoA-producing enzyme in H. pylori,
Acetate kinase	ackA	COG0280	Disrupted by a frameshift	
Phospho- transacetylase	<i>p</i> ta	COG0278	Disrupted by frameshifts	
Enzymes of purine	purF	COG0034	Missing	De novo purine biosynthesis is absent in H. pylori, and it
biosynthesis	purD	COG0151	Inactivated by mutations	has to obtain purines from the host. HP1185 appears to be the best candidate for the purine permease, as it is the only
	ригМ	COG0299	Missing	H. pylori protein, similar to E. coli PurP.
	purT	COG0027	Missing	
	purL_1	COG0046	Missing	
	purL_2	COG0047	Missing	On the other hand, H. pylori encodes the enzymes for AMF
•	purM	COG0150	Missing	and GMP synthesis from IMP and their interconversion.
	purK	COG0026	Missing	Therefore, it can survive on any of these purines.
	purE	COG0041	Missing	•
	purC	COG0152	Missing	
	purH	COG0138	Missing	
	purA	COG0104	Present	
	purB	COG0015	Present	
	guaB	COG0516	Present	
	guaA_1	COG0518	Present	
	guaA_2	COG0519	Present	

sive duplication with divergence in the eukaryotes. The expansion of this domain into a number of eukaryotic proteins involved in cell-cycle control [50",51] may have been critical for the very establishment of these systems.

With the current acceleration in protein structure determination [22,24], a superfamily identified by sequence comparison more and more frequently extends to include proteins with known 3D structure and/or well-characterized catalytic mechanism (Table 3). Such findings are sometimes most illuminating as they immediately result in the prediction of the structural fold, the structure of the active center, and possibly also the catalytic mechanism for a wide variety of diverse proteins comprising the superfamily. This is illustrated by the recent prediction of the structure and the catalytic amino acid residues for P-ATPases, which remained elusive in spite of a long history of studies, on the basis of the sequence motifs shared with haloacid dehalogenases [52*].

Assignment of the gene products to structural folds and families with maximal attainable precision is arguably one of the foremost tasks of genome analysis after the sequencing phase. The number of structures that have been determined experimentally is negligible for almost all genomes, with the exception of E. coli (where it is still rather a small fraction) (Table 1). A database search with a deliberately conservative similarity cut-off already increases the fraction of proteins for which a confident structure prediction is possible to 10-25% [53'] (Table 1). Secondary structure-based threading allows

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Superfamily	Enzymes with known 3D structures (PDB codes)	Enzymes with newly predicted properties	Representatives in complete genomes*	References
BRCT (conserved domain in cell cycle checkpoint proteins)	None	DNA polymerase subunit DPB11, terminal deoxynucleotidyltransferase, DNA figases III and IV, poly(ADP-ribose) polymerase	e-1, h-1, u-1, b-1, g-1, p-1, l-1, c-2, m-0, t-0, f-0, y-9, w-8	[20]
Urease-related metaf-dependent hydrolases	Urease (2kauC), phosphotriesterase (1pta), adenosine deaminase (1fkx)	AMP deaminase, adenine deaminase, cytosine deaminase, hydantoinase, dihydroorotase, allantoinase, aminoacyłase, imidazolonepropionase, arylphosphatase, chlorohydrolase, formylmethanofuran dehydrogenase	e-13, h-4, u-2, b-6, g-1, p-1, l-1, c-3, m-9, t-10, f-7, y-6, w-9	[52]
Acid phosphatases	Vanadium-containing chloroperoxidase (1 vnc)	Phosphatidic acid phosphatase, phosphatidylglycerol phosphatase, diacylglycerol pyrophosphate phosphatase, glucose-6-phosphatase	e-4, h·1, u-3, b-2, g-0, p-0, l-0, c-1, m-3, t-1, f-0, y-7, w-4	[58,57]
ATP-grasp (ATP-dependent C-N and C-S ligases)	Glutathione synthetase (1gsh), D-ala-D-ala ligase (2din), biotin carboxylase (1bnc), carbamoy! phosphate synthase (1jdb), succinyl-CoA	Phosphoribosylamine-glycine ligaso, phosphoribosylgłycinamide formyltransferase, phosphoribosylaminoimidazole carboxylase, tubulin-tyrosine ligase, protein S6-glutamate ligase (RimK), malate thiokinase, ATP-citrate lyase	e-10, h-5,u-4, b-12, g-2,p-2, f-1, c-7, m-9, t-8, f-11, y-10, w-15	[28]
HAD (phosphatases and other hydrolases)	L·Haloacid dehalogenase (1 jud)	Phosphoserine phosphatase, phosphoglycolate phosphatase, histidinol phosphatase, glycerol-3-phosphatase, sucrose phosphate synthase, phosphomannomutase, P-type cation-transport ATPases	e-10, h-3, u-1, b-11, g-4, p-5, l-2, c-8, m-3, t-4, f-3, y-9, w-6	[52]
· DHH (hydrolases)	None	Exopolyphosphatase, 5'-3' exonuclease	e-1, h-1, u-4, b-6, g-2, p-2, l-2, c-3, m-6, t-3, f-7, y-1, w-0	[69]
Alkaline phosphatase- related metal- dependent hydrolases	Alkaline phosphatase (talk), N-acetylgalactosamine 4-sulfatase (tfsu), cerebroside sulfatase(tauk)	Phosphopentomutase, 2,3-bisphosphoglycerate-independent phosphoglycerate mutase, streptomycin-6-phosphatase, phosphonoacetate hydrolase, phosphoglycerol transferase, nucleotide pyrophosphatase, steroid suffatase,	e-15, h-5, u-4, b-8, g-1, p-1, l-0, c-1, m-2, t-2, f-3, y-6, w-18	[40,60]

^{*}Identified by PSI-BLAST [47**] searches of the complete genomes using the conserved motif(s) for each superfamily as a query. Organism abbreviations are as in Figure 1: e, E. coli; h, H. influenzae; u, H. pylori; b, B. subtilis; g, M. genitalium; p, M. pneumoniae; l, B. burgdorferi; c, Synechocystis sp.; m, M. jannaschii; t, M. thermoautotrophicum; t, A. fulgidus; y, S. cerevisiae; w, C. elegans.

another relatively small but notable increase in the predictive power [54] (Table 1). It appears, however, that at this time, the most realistic way to further structure prediction at genome scale is to perform a complete analysis of protein superfamilies as exemplified in Table 3.

Perspective

As far as prokaryotic genomes are concerned, we have already entered the post-genomic era. While surprises certainly wait ahead, there is little doubt that the major protein families are already known or can be deciphered from the available sequences. We have recently seen major progress in methods and procedures for advanced sequence analysis, and a lot of valuable information has been extracted from the genomes. We believe, however, that a major focused effort in genome comparison is still required in order to construct a proper classification of protein families and superfamilies and systematically apply it to the goals of structural and functional prediction. Such an effort will have the potential of creating a basis for a rationally designed, decisive onslaught on structure determination and experimental identification of gene functions using computer predictions as a guide. Hopefully, this research program turns out to be both realistic and efficient.

References and recommended reading

Papers of particular interest, published within the annual period of review, have been highlighted as:

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- of outstanding interest
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The completion of the genome sequence of E. coli, one of the classic objects of molecular biology and genetics, certainly has a symbolic significance. More importantly, the enormous amount of information available regarding E. coli gene functions can now be used to full potential for inferring functions of homologs in other species. However, the functions of about one half of the E. coli genes have not been determined experimentally, and so there is still a lot to learn about E. coli itself.

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Nature 1997, 388:539-547.

The genome sequence of this bacterium is of special interest from several points of view. The genome analysis will have important practical implications as H. pylori is the causative agent of peptic ulcers and is believed to infect up to half of the human population. H. pylori thrives in a highly acidic environment (pH 2-3); deciphering the mechanisms of acid tolerance from the genome sequence is a most interesting task. Furthermore, H. pylori represents an early branching of the proteobacterial lineage, and the comparison of its genome with those of other Proteobacteria such as E. coli and Haemophilus influenzae will shed light on the evolution of cellular functions in bacteria and mitochondria.

Smith DR, Doucette-Stamm LA, Deloughery C, Lee H, Dubois J, Aldredge T, Bashirzadeh R, Blakely D, Cook R, Gilbert K et al.: Complete genome sequence of Methanobacterium thermoautotrophicum deltaH: functional analysis and comparative genomics. J Bacteriol 1997, 179:7135-7155.

This second genome of a methanogenic archeon to be sequenced, after Methanococcus jannaschii, is of major importance in corroborating trends revealed by the M. jannaschii genome analysis [4,34]. Like M. jannaschii, there is a sharp divide between the majority of the genes, which appear to have bacterial origin, and a minority (primarily encoding proteins involved in genome replication and expression) of 'eukaryotic' genes. Some other unusual aspects of the M. jannaschii genome, however, did not recur in M. thermoautrophicum. For example, unlike M. jannaschii, M. thermoautrophicum encodes a typical set of molecular chaperones such as DnaK and DnaJ and does not encode a unique ATPase family found in M. jannaschii.

10. Klenk HP, Clayton RA, Tomb JF, White O, Nelson KE, Ketchum KA, Dodson RJ, Gwinn M, Hickey EK, Peterson JD et al.: The complete genome sequence of the hyperthermophilic, sulphate-reducing archaeon Archaeoglobus fulgidus. Nature 1997, 390:364-370.

The first sequence of a non-methanogenic archeon, and the third complete archeal genome altogether. With 2436 genes, the A. fulgidus genome is considerably larger than those of M. jannaschii and M. thermoautrophicum, in part due to more extensive duplication in some of the gene families. Unlike M. jannaschii and M. thermoautrophicum, A. fulgidus does not seem to encode any inteins. With three genome sequences available, there is for the first time an opportunity for an informative comparative analysis of archeal genomes. Definitive work in this area remains to be done, but it is already clear that the three genomes generally are highly coherent, and also that there are many mysterious conserved families, creating a challenge for further research, both theoretical and experimental.

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The second classic bacterial model, after E. coli, and also the second largest bacterial genome sequenced so far (4100 genes compared with 4288 genes in E. coli). With B. subtilis adequately representing the Grampositive lineage (only the minimal genomes of Mycoplasma had been available before), we may now have a sampling of the great majority of bacterial gene families. In addition to its value for comparative analysis, B. subtilis is most interesting and important in its own right, given, for example the large number of genes in its genome that encode enzymes of secondary metabolite synthesis.

12. Fraser CM, Casjens S, Huang WM, Sutton GG, Clayton R, Lathigra •• R, White O, Ketchum KA, Dodson R, Hickey EK et al.: Genomic sequence of a Lyme disease spirochaete, Borrella burgdorferi. Nature 1997, 390:580-586.

The first genome representing yet another major division of bacteria, the spirochetes. The genome has a number of unique features, above all a linear chromosome unusual in the bacterial world, and at least 17 linear and circular plasmids that contain about 30% of the genes. Most of the plasmid-borne genes remain quite mysterious, at least after the initial genome analysis.

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A detailed comparison of the first available archeal genome (M. jannaschii) with bacterial genomes produced a number of novel functional predictions and led to the conclusion that the majority of archeal genes most probably have a bacterial origin. Furthermore, generalizations started to emerge, including the nearly constant fraction of genes containing ancient conserved regions - about 70% in all genomes - and the same major superfamilies of paralogs.

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Comparative analysis of the proteins encoded in seven complete genomes from five major phylogenetic lineages and elucidation of consistent patterns of sequence similarities resulted in the delineation of 720 clusters of orthologous groups (COGs). Each COG consists of individual orthologous proteins or orthologous sets of paralogs from at least three lineages. Orthologs typically have the same function, allowing transfer of functional information from one member to an entire COG. This automatically makes possible a number of functional predictions, especially for poorty characterized genomes. The evolving system of COGs comprises a

framework for functional and evolutionary genome analysis; it is accessible through the World Wide Web (http://ncbi.nlm.nih.gov/COG).

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A major revamp of BLAST, which is definitely the most popular current method for database search. The key innovations are: first, the program now makes gapped alignments, with appropriately modified statistics, which results in significant increase of sensitivity; and second, the associated program PSI (Position-Specific Iterating)-BLAST makes a position-specific weight matrix (profile) out of the first pass results and iterates searches with this profile until no new sequences with similarity scores above a defined cut-off are detected. This appears to be the most powerful existing method for detection of subtle similarities between protein sequences and delineation of protein superfamilies.

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Sequence analysis of the proteins encoded by 70 positionally cloned human disease genes showed that most of them have orthologs with the same domain architecture in the nematode, but domain rearrangements are prevalent in yeast and bacterial homologs. This is one of the first demonstrations of the utility of PSI-BLAST for the defineation of large protein superfamilies. In particular, this method was used for the identification of a conserved ATPase domain present in the repair protein MutL (one of the colon cancer gene products in humans), histidine kinases, molecular chaperones of the HSP90 family and type II DNA topoisomerases; the 3D structure for the latter was already available, defining the fold for the whole superfamily.

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A complete description of the BRCT domain that had been originally found in BRCA1 protein and several other proteins implicated in cell cycle checkpoint. In this work, the superfamily has been extended to include a distinct version of the BRCT domain detected in bacterial DNA ligases, the large subunits of eukaryotic replication factor C, and poly(ADP-ribose) polymerases. The expansion of the BRCT domain in eukaryotes may be one of the key events in the evolution of cell-cycle control.

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This paper describes a very convenient Worldwide Web site compiling results of automatic analysis of all available complete genomes. The Pedant WWW site (http://pedant.mips.biochem.mpg.de/frishman/pedant.html) is arguably one of the best entry points to comparative genomics but it has to be kept in mind that it is only the first level, crude analysis that is presented here.

 Fischer D, Eisenberg D: Assigning folds to the proteins encoded by
 the genome of Mycoplasma genitalium. Proc Natl Acad Sci USA 1997, 94:11929-11934.

One of the first systematic attempts to predict the 3D structures of proteins starting from a complete genome. The utility of sequence-structure threading is demonstrated but it also becomes clear that such methods at best result in a rather small, incremental improvement over state-of-the-art sequence comparisons. Although the fraction of the proteins with a

predictable fold is only 22% of the gene products, the authors predict by extrapolation that it should be possible to assign folds to most soluble proteins within a decade.

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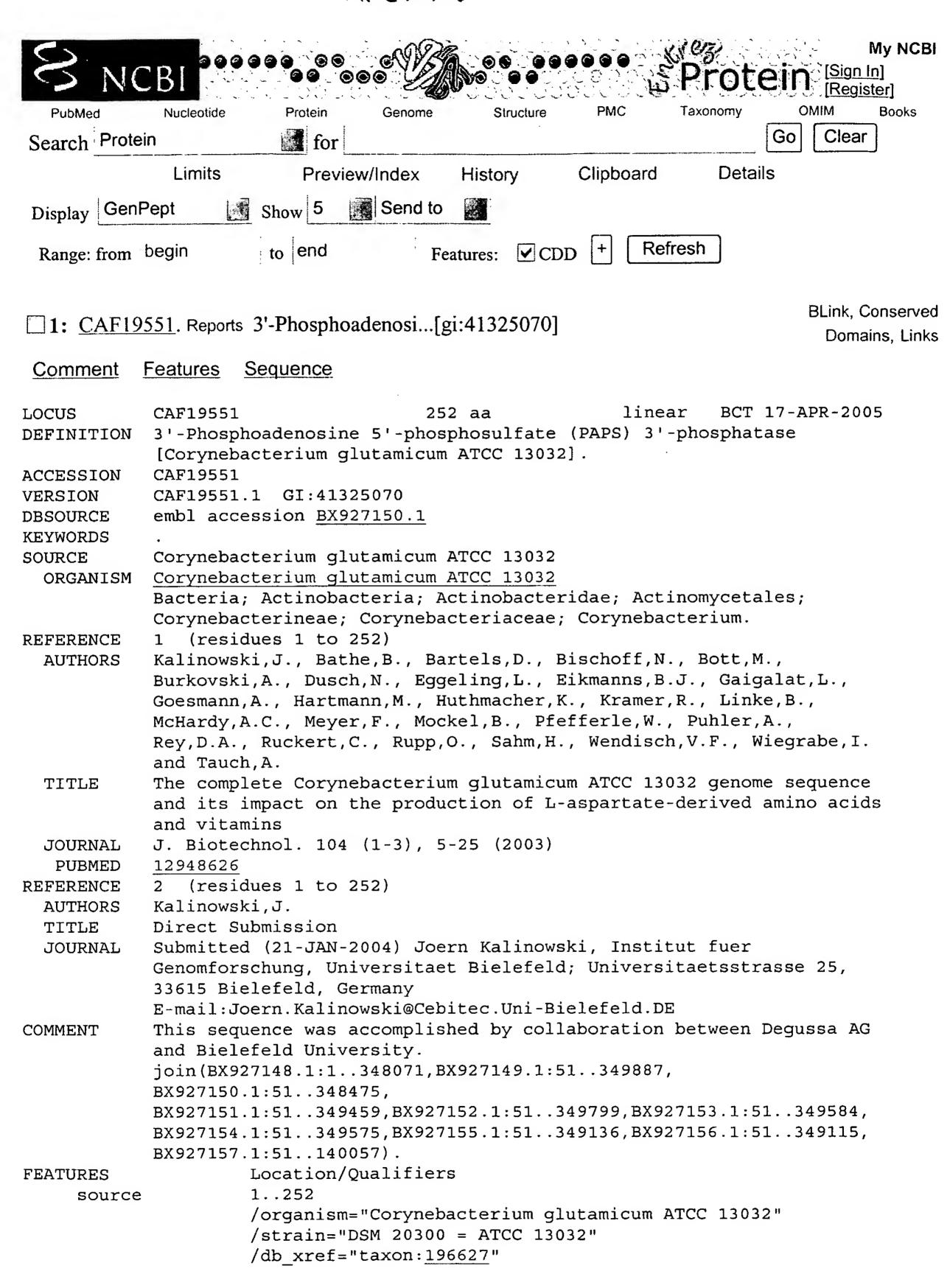
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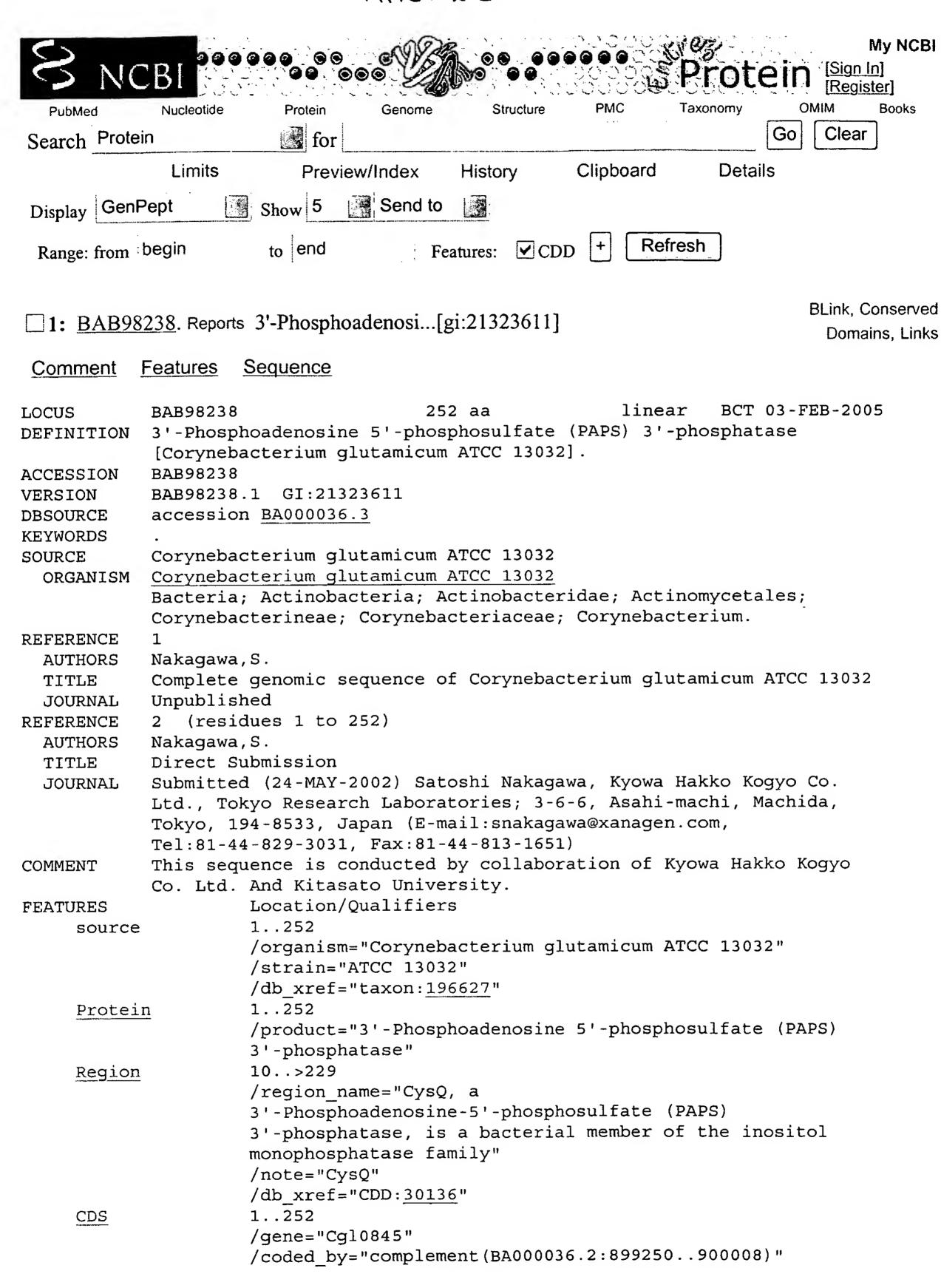
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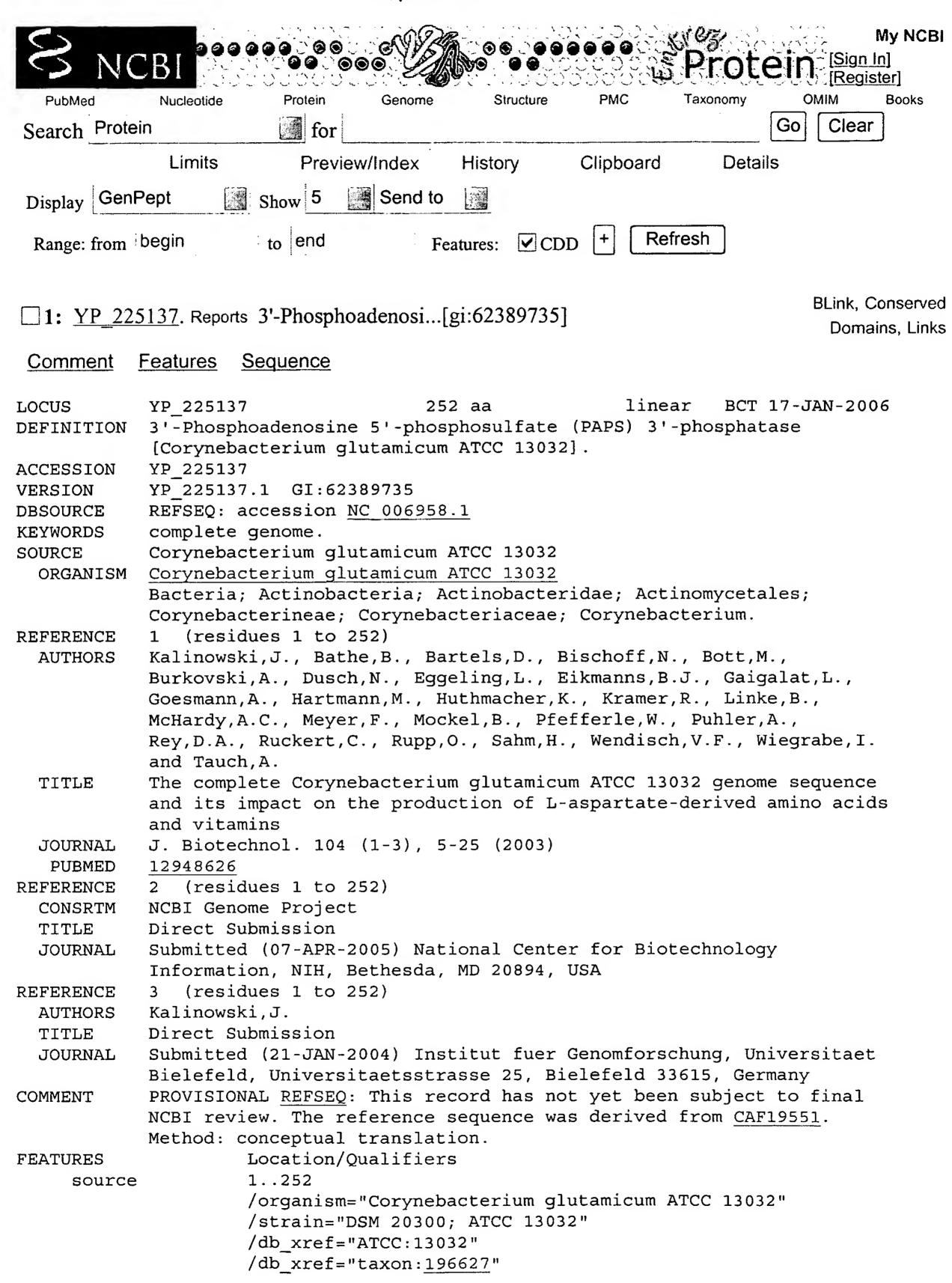
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[ISREC-Server] Date: Mon Jun 26 19:28:01 Europe/Zurich 2006

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CAF195	MTAQIDDSILTHRLA			NLGDAGDELA	AOSWIARVLE	OHRPNDG
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Back to ISREC bioinformatics group home page

APPENDIX 4

lalign output for SEQ ID NO:6 vs. BAB98238

[ISREC-Server] Date: Mon Jun 26 19:35:51 Europe/Zurich 2006

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SEO ID NO:6
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BAB98238
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                                                 50
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lalign output for SEQ ID NO:6 vs. YP_225137

[ISREC-Server] Date: Mon Jun 26 19:39:50 Europe/Zurich 2006

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cysQ, a Gene Needed for Cysteine Synthesis in Escherichia coli K-12 Only during Aerobic Growth

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The initial steps in assimilation of sulfate during cysteine biosynthesis entail sulfate uptake and sulfate activation by formation of adenosine 5'-phosphosulfate, conversion to 3'-phosphoadenosine 5'-phosphosulfate, and reduction to sulfite. Mutations in a previously uncharacterized Escherichia coli gene, cysQ, which resulted in a requirement for sulfite or cysteine, were obtained by in vivo insertion of transposons Tn5tac1 and Tn5supF and by in vitro insertion of resistance gene cassettes. cysQ is at chromosomal position 95.7 min (kb 4517 to 4518) and is transcribed divergently from the adjacent cpdB gene. A Tn5tac1 insertion just inside the 3' end of cysQ, with its isopropyl-β-D-thiogalactopyranoside-inducible tac promoter pointed toward the cysQ promoter, resulted in auxotrophy only when isopropyl-β-p-thiogalactopyranoside was present; this conditional phenotype was ascribed to collision between converging RNA polymerases or interaction between complementary antisense and cysQ mRNAs. The auxotrophy caused by cysQ null mutations was leaky in some but not all E. coli strains and could be compensated by mutations in unlinked genes. cysQ mutants were prototrophic during anaerobic growth. Mutations in cysQ did not affect the rate of sulfate uptake or the activities of ATP sulfurylase and its protein activator, which together catalyze adenosine 5'-phosphosulfate synthesis. Some mutations that compensated for cysQ null alleles resulted in sulfate transport defects. cysQ is identical to a gene called amtA, which had been thought to be needed for ammonium transport. Computer analyses, detailed elsewhere, revealed significant amino acid sequence homology between cysQ and suhB of E. coli and the gene for mammalian inositol monophosphatase. Previous work had suggested that 3'-phosphoadenoside 5'-phosphosulfate is toxic if allowed to accumulate, and we propose that CysQ helps control the pool of 3'-phosphoadenoside 5'-phosphosulfate, or its use in sulfite synthesis.

The cysteine biosynthetic pathway (Fig. 1), a principal route of sulfur assimilation, involves more than 15 genes in at least five chromosomal regions in Escherichia coli and Salmonella typhimurium. It has been studied since the early days of physiological genetics in order to elucidate the roles of the individual genes, the control of their expression, and how the flow of metabolic intermediates is regulated (for a review, see reference 25). The transcription of most cys genes is positively controlled by the protein product of cysB and its coinducer, O-acetyl serine (also a cysteine precursor), during aerobic growth; transcription is repressed by sulfide, which is generated by reversal of the final biosynthetic step (Fig. 1). CysB seems not to be needed during anaerobic growth (3). The cysQ gene described here is also needed only during aerobic growth. It is inferred to act before sulfite formation, and hence this early part of the cysteine pathway is reviewed briefly below.

The initial step, sulfate uptake, is mediated by a permease encoded by the cysT, cysW, and cysA genes, which, along with cysP, constitute one operon (49). CysP protein is needed for maximal thiosulfate and sulfate binding, but it is probably not part of the permease, and its role in cysteine

biosynthesis is unclear (19). A cysZ gene, about 10 kb from the cysPTWA operon, may also be needed for sulfate uptake (41). Intracellular sulfate is activated via synthesis of adenosine 5'-phosphosulfate (APS) by ATP sulfurylase, which is encoded by cysD and cysN (34). This activation step is complex, in that the rate of APS formation is greatly enhanced both by a protein activator (31, 32) and by GTP hydrolysis (33). APS is converted to 3'-phosphoadenosine 5'-phosphosulfate (PAPS) by APS kinase, encoded by cysC. This step is thought to not require cofactors, because APS kinase activity does not change during enzyme purification (46). Sulfite is generated from PAPS in a complex reaction involving transfer and reduction of its sulfuryl moiety. This reaction is catalyzed by the cysH gene product, PAPS sulfotransferase, and involves a thioredoxin- or glutaredoxin-bound intermediate (51, 52).

Strains with mutations in cysH or in both trxA and grx (encoding thioredoxin and glutaredoxin, respectively) grow poorly. The poor growth can be corrected by additional mutations in cysC (APS kinase) or genes for earlier steps in the pathway (16, 45a), a result indicating that PAPS or one of its derivatives is toxic if allowed to accumulate. We find this result interesting in the context of understanding mechanisms by which organisms cope with the many metabolic intermediates that are both essential for healthy growth and potentially deleterious. Other studies have shown that the activities of ATP sulfurylase and APS kinase decrease rapidly when growth is slowed (25, 26). Such instability could help modulate metabolite flow through this pathway and would be more sensitive to decreased need for PAPS

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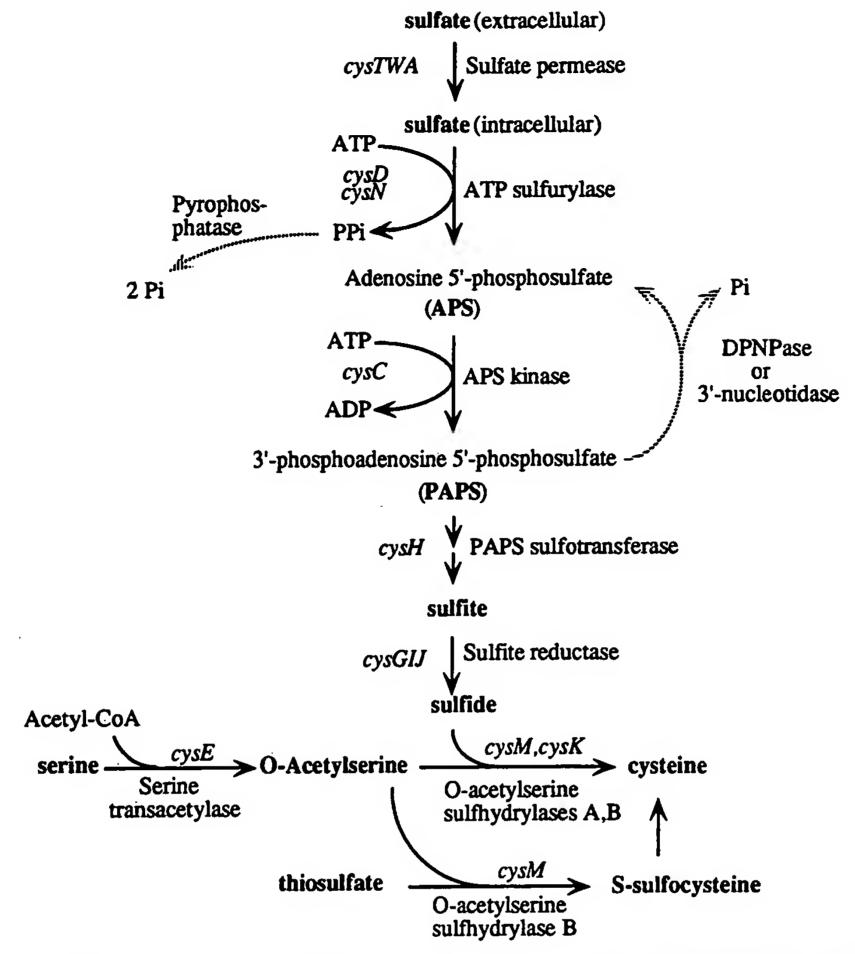


FIG. 1. Pathway of cysteine biosynthesis in E. coli (modified from data in references 25, 34, and 49).

than any transcriptional regulation. The dependence of APS synthesis (and thereby PAPS synthesis) on the ATP sulfury-lase activator and on the local concentration of GTP (32, 33) might also help regulate PAPS levels.

Mutations in the cysQ gene described here result in a requirement for cysteine or sulfite that is expressed only during aerobic growth and that is leaky in many but not all laboratory strains of $E.\ coli$. Our studies suggest that CysQ may help control the levels of PAPS, its localization, or its use in sulfite synthesis.

MATERIALS AND METHODS

Strains, media, and general methods. The bacterial strains and plasmids used in this study are listed in Table 1. Bacteria were grown in LN broth (5) or Vogel-Bonner glucoseminimal salts medium (54). An M9-based minimal salts medium, with ammonium acetate in place of ammonium chloride (21), was used where indicated. Solid media contained 1.5% Difco Bacto-Agar. Antibiotics were used at the following concentrations: ampicillin, 250 μg/ml; kanamycin, 60 μg/ml; tetracycline, 12 μg/ml; streptomycin, 100 μg/ml; and chloramphenicol, 20 μg/ml. Isopropyl-β-D-thiogalacto-

pyranoside (IPTG) was used at 0.5 mM. Amino acids were added at 50 μg/ml except for glycine, which was added at 200 μg/ml. Standard procedures were used for bacterial growth, characterization of auxotrophy and conjugation, DNA preparation, restriction endonuclease digestion, DNA electrophoresis, recombinant DNA cloning, and transformation (12, 47). All enzymes were obtained from commercial sources (Life Technologies, Inc., Stratagene, New England BioLabs, or Boehringer Mannheim) and used as directed. Anaerobic (H₂-CO₂ atmosphere) conditions were obtained by using BBL GasPak Anaerobic jars and the BBL GasPak plus system (Becton Dickinson and Co.).

Assays. The activity of 2',3'-cyclic phosphodiesterase (encoded by the cpdB gene) was measured as the release of inorganic phosphate from cyclic UMP (4). Sulfate uptake was measured as depletion of $^{35}SO_4$ added to the medium with cells grown with djenkolic acid as the sulfur source and concentrated from the exponential phase (19). ATP sufury-lase was measured as incorporation of $^{35}SO_4$ into PAPS, detected by thin-layer chromatography with dialyzed extracts of cells grown with sulfite as the sulfur source and induced with O-acetyl-1-serine (34). The level of the activator of ATP sulfurylase was also measured as PAPS synthesis

TABLE 1. Bacterial strains, phage, and plasmids

Strain, phage, or plasmid	Description or genotype	Source or reference
E. coli		
AJ2653 ^a	ET8000 amtA (cysQ)::Tn10	21
BW6458	proC::Tn5 zje::Tn10-BJW43 metB1 relA1	B. Wanner
CAG5052	Hfr btuB3191::Tn10, transfer counterclockwise from 7 min	48
DB747	W3350 gal rpsL sup ⁰ (strain 594 of reference 7)	Laboratory collection
DB1434	DB747 (λplac5 c1857 Sam7)	28
DB4496	MC1061 dam::Tn9 (p3)(pBRG1310)	43
DB5463	HfrH lacZ(Am) trp(Am) sup ⁰	D. Botstein (DB6128)
DB5508	recD (p3)	44
DB5659	DB747 cysQ::kan	λcysQ::kan transduction
DB6302 ^a	MG1655 amtA (cysQ)::Tn10	P1 transduction from AJ2653
DB6316	MG1655 cysQ::kan	$\lambda cysQ::kan$ transduction
DB6908	ET8000 cysQ::kan	$\lambda cysQ::kan$ transduction
DB6913	TG1 cysQ::kan	$\lambda cysQ::kan$ transduction
DB6935	DB5508 cysQ::Tn5supF	This study
DB7101	TG1 $\Delta(cysD-N)$::kan	$\lambda \Delta (cysD-N)$::kan transduction
DBan41	594 with cysQ::Tn5tacl	Tn5tac1 transposition
DBan41TR	DBan41 with Δ (srl-recA)306::Tn10	P1 transduction from JC1289
DK21	sup ⁰ dnaB(Am) 266 (λimm ²¹ -ban _{P1})	29
ET8000	rbs lacZ::IS1 gyrA hutC _k	37
JC1289	$\Delta(srl-recA)306$ linked to Tn10	11
MC1061	F⁻araD139∆(ara-leu)7697 ∆lacX74 galU galK hsdR hsdM rpsL	8
MG1655	F ⁻ prototroph	17
TG1	F' proAB ⁺ traD36 lacI ^q lacZ Δ M15 supE hsd Δ 5 thi Δ (lac-proAB)	47
BW6164	HfrRA2 thr::Tn10, clockwise transfer from 88 min	55
Phages		
M13mp18	Cloning vector	38
λ^+	λ wild type	Laboratory collection
λ::Tn5tacl	λTn5tac1 b221 c1857 Oam29 Pam80	9
λ419	cysA ⁺ (5F7 of reference 24)	23
λ656	$cysQ^+$ (5B5 of reference 24)	23
$\lambda \Delta (cysD-N)$::kan	Derivative of λ652 (6C8 of reference 24)	28
λcysQ::kan	Derivative of λ656	28
$\lambda 656cysQ::Tn5supF$		This study
Plclr		Laboratory collection
Plasmids	A T . O TO C . A /W D . O+	
pBRGan101	Amp ^r , cysQ::Tn5tacl (Kan ^r) cysQ ⁺	pBR322 (Sall fragment from DBan41)
pBRGan102	Amp ^r , $cysQ$::Tn5tac1 (Kan ^r) $cysQ^+$ $cpdB^+$	pBR322 (Clal fragment from DBan41)
pBRGan103	Amp ^r , $cysQ^+$ $cpdB^+$ Clal::Tn5tacl $\Delta(0-52)$	pBRGan102, small ClaI deletion (Kan ^r)
pBRGan104	Amp^r , $cpdB^+$	pBRGan102, EcoRI deletion
pBRGan110	Amp^r , $cysQ^+$	pBR322 (SalI fragment of DB747)
pBRGan111	Amp ^r , $cysQ^+$	pAN110, partial MspI digestion
pBRGan111-1 to pBRGan111-5	Amp ^r , cysQ::cat	cat of pCM4 ligate into partial Sau3A of pBRGan111 (Fig. 2 and 4)
pcysQ::kan	Amp ^r , cysQ::kan	kan of pUC4K into EcoRI site of pBRGan111
pCM4		10
pBR322	Amp ^r Tet ^r	6
p3	Kan ^r amp(Am) tet(Am)	29
pBRG1310	Tn5supF donor	43

^a The amtA gene is identical to cysQ, as detailed in the text.

in reactions containing purified ATP sulfurylase and dialyzed cell extracts as the source of ATP sulfurylase activator (31, 32).

Genetic manipulation and analysis. Standard methods were used for (i) mutagenesis of $E.\ coli$ with Tn5tac1 with phage $\lambda::Tn5tac1$ b221 c1857 Oam29 Pam80 as a transposon donor (9) and (ii) Hfr conjugation and P1 generalized transduction (48). Cysteine-requiring bacteria were tested for sensitivity to azaserine and to chromate by spotting dilutions of these agents on lawns of 10^7 bacteria spread on minimal glucose

agar supplemented with cysteine, djenkolic acid, or glutathione and IPTG, as appropriate, and by growth in liquid cultures with progressive twofold differences in the concentrations of these agents.

To insert a transcription reporter into the cysQ gene, the DNA of cysQ plasmid pBRGan111 was partially digested with Sau3A, and full-length linear DNA was isolated after electrophoresis in low-melting-point agarose and ligated to the BamHI cat fragment from plasmid pCM-4 (10). The

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religated DNA was used to transform the cysQ::Tn5tacl strain DBan41, and plasmids that did not complement its cysteine auxotrophy were identified and characterized.

To generate a cysQ::Tn5supF insertion mutant, Tn5supF was transposed from the donor plasmid in strain DB4496 (43) to $cysO^+$ phage $\lambda656$ (23, 24), and insertion-containing phage were selected by plaque formation on the dnaB amber strain DK21 (29, 43). Haploid Tn5supF-containing bacterial recombinants were obtained by infecting strain DB5508 (which contains amber mutant alleles of amp and tet genes) and selecting Sup⁺ transductants by their resistance to ampicillin or tetracycline (44). Sup⁺ transductants were screened for auxotrophy. To generate a cysQ::kan mutant, an EcoRI kan cassette from plasmid pUC4K was ligated into the EcoRI site in cysQ of pBRGan111. This allele was recombined into λ656 by infecting cells carrying the pBRGan111-cysQ::kan plasmid and selecting phage carrying the cysQ::kan allele by transduction of DB1434. λcysQ::kan phage recovered from the lysogen were used to transduce nonlysogens and thereby obtain haploid cysQ::kan bacteria (28).

Cys⁺ revertants of cysQ mutant strains were obtained by growing young single-colony isolates in 2 ml of LN broth to stationary phase, washing the cells twice with 10 mM MgSO₄, plating aliquots on minimal (cysteine-free) medium, and incubating for 2 days at 37°C. Reversion frequencies were measured by using several cultures from different single colonies to avoid jackpots.

DNA sequence analysis. A 1-kb segment containing the cysQ gene was sequenced by the Sanger dideoxynucleotide-chain termination method with Sequenase (U.S. Biochemical, Cleveland, Ohio) and single- and double-stranded DNA templates (27). Primer binding sites were provided by insertions of transposons Tn5tacl and Tn5supF in phage λ 656, by the promoterless cat gene in plasmid pBRGan111 DNAs, and by a universal primer binding site in M13mp18 (38) (for sequencing an EcoRI-PstI fragment containing the 3' end of cysQ).

The oligonucleotides used as sequencing primers are as follows: (i) 5' CTCCATTTTAGCTTCCTTAGCTCC, positions 40 through 17 at the 5' end of the cat gene cassette; (ii) 5' TGTCAAAACATGAGAATTCCTCCCG, positions 43 through 20 near the I end of Tn5tacl; (iii) 5' GGAAACAGA ATTCCCGGGGGATCCCC, positions 4549 through 4573 near the O end of Tn5tacl; (iv) 5' TAGGATCCCCTACTTGT GTA, positions 30 through 11 near the O end of Tn5supF; (v) 5' TAGGATCCCGAGATCTGATC, positions 236 through 255 near the I end of Tn5supF; (vi) 5' GAGCGGCC AAAGGGAGCAGAC, positions 139 through 159 (middle primer) within Tn5supF with its 3' end toward the I end; (vii) 5' GTAAAACGACGGCCAGT, the universal M13 sequencing primer.

Nucleotide sequence accession number. The nucleotide sequence of the cysQ gene shown in Fig. 4 has been deposited with GenBank under accession number M80795.

RESULTS

Initial detection and characterization of cysQ. The prototrophic strain E. coli DB747 was mutagenized with Tn5tacl, a transposon with an outward-facing tac promoter that is regulated by the lac repressor and IPTG (9). A conditional mutant that required cysteine for growth on minimal medium containing IPTG, but not on medium lacking IPTG, was isolated and named DBan41. Early characterizations of this strain revealed two other novel features. It did not require cysteine for normal growth in an anaerobic

atmosphere. In addition, it formed slow-growing colonies on cysteine-free medium containing IPTG (after 2 to 3 days, instead of 16 h in the case of its Cys⁺ parent). Cells in these colonies exhibited the same slow-growth phenotype, which indicated that the mutation was leaky, not highly revertible. The addition of IPTG to DBan41 in cysteine-free liquid medium lengthened the cell doubling time from about 80 min to 280 min.

The cysteine requirement of DBan41 was satisfied by sulfite at 0.3 mM, which indicated a defect in the sulfate assimilation branch of the pathway (Fig. 1). The strain was as sensitive to chromate (MIC, 50 to 100 µM) as its wild-type parent and also grew on low concentrations of thiosulfate (1 to 2 mM). Sulfate uptake mutants are chromate resistant, and many are deficient in thiosulfate uptake (14, 40); therefore this mutation seemed to affect a step leading to sulfite that follows sulfate uptake (Fig. 1).

The mutation was mapped by genetic and molecular methods. (i) The cys^+ allele was transferred efficiently by Hfr strains BW6164 and CAG5052 to DBan41, which placed the mutation in the 88- to 07-min interval of the $E.\ coli$ chromosome, far from other known cysteine biosynthetic genes (1). (ii) The cys^+ allele was cotransduced by phage P1 at a frequency of 1% with zje::Tn10, an insertion at 94 to 95 min (strain BW6458). (iii) The cys^+ allele was also efficiently transduced by $\lambda656$ (23, 24), a λ phage clone that carries the segment of the $E.\ coli$ chromosome from kb \sim 4511 to kb \sim 4525 (near 96 min). The mutant allele was recessive to the wild type in partial diploids, which were formed by $\lambda656$ infection of a λ^+ lysogenic derivative of DBan41 (44), as well as in strains carrying the wild-type allele in multicopy plasmids.

More refined map information came from molecular cloning (Fig. 2). (i) Kan^r plasmids obtained by cloning SalI- or ClaI-digested DBan41 DNA in pBR322 contained an 18-kb SalI fragment (pBRGan101) or an overlapping 14-kb ClaI fragment (pBRGan102), respectively. (ii) A plasmid obtained by cloning SalI-digested wild-type E. coli DNA (pBRGan110) that complemented the cysQ::Tn5tacl allele contained a 14-kb fragment whose restriction map matched that of the chromosome adjacent to the Tn5tacl insertion. (iii) A deletion plasmid that retained only 1.8 kb of chromosomal sequence but retained Cys⁺ complementing activity was generated by partial MspI digestion of pBRGan110 DNA (pBRGan111). Comparisons of the restriction digest patterns of these clones with the known restriction map of the chromosomal region near 96 min (24, 36) indicated that Tn5tacl was at kb 4517, about 900 bp upstream of the cpdB gene. Transcription from the tac promoter in Tn5tac1 was toward cpdB (clockwise).

Cys⁻ phenotype not caused by cpdB overexpression. The CpdB protein has a 3'-nucleotidase activity that can degrade PAPS to APS in vitro (4). Although CpdB protein seems to be primarily periplasmic, findings of cytoplasmic inhibitors for other periplasmic nucleotidases (36) suggested models in which CpdB also acted intracellularly. Thus, in principle, transcription from the tac promoter might cause a cysteine requirement by increasing cpdB expression. Alternatively, it might alter the expression of an unknown gene next to cpdB. Three findings eliminated the simple CpdB-based model of cysteine auxotrophy: (i) the CpdB level in strain DBan41 was increased less than 2-fold by IPTG (data not shown); (ii) the multicopy cpdB⁺ plasmid (pBRGan104) did not cause a cysteine requirement, although it did result in 10-fold higher CpdB activity (data not shown); and (iii) the cysteine requirement was complemented by plasmid pBRGan102.

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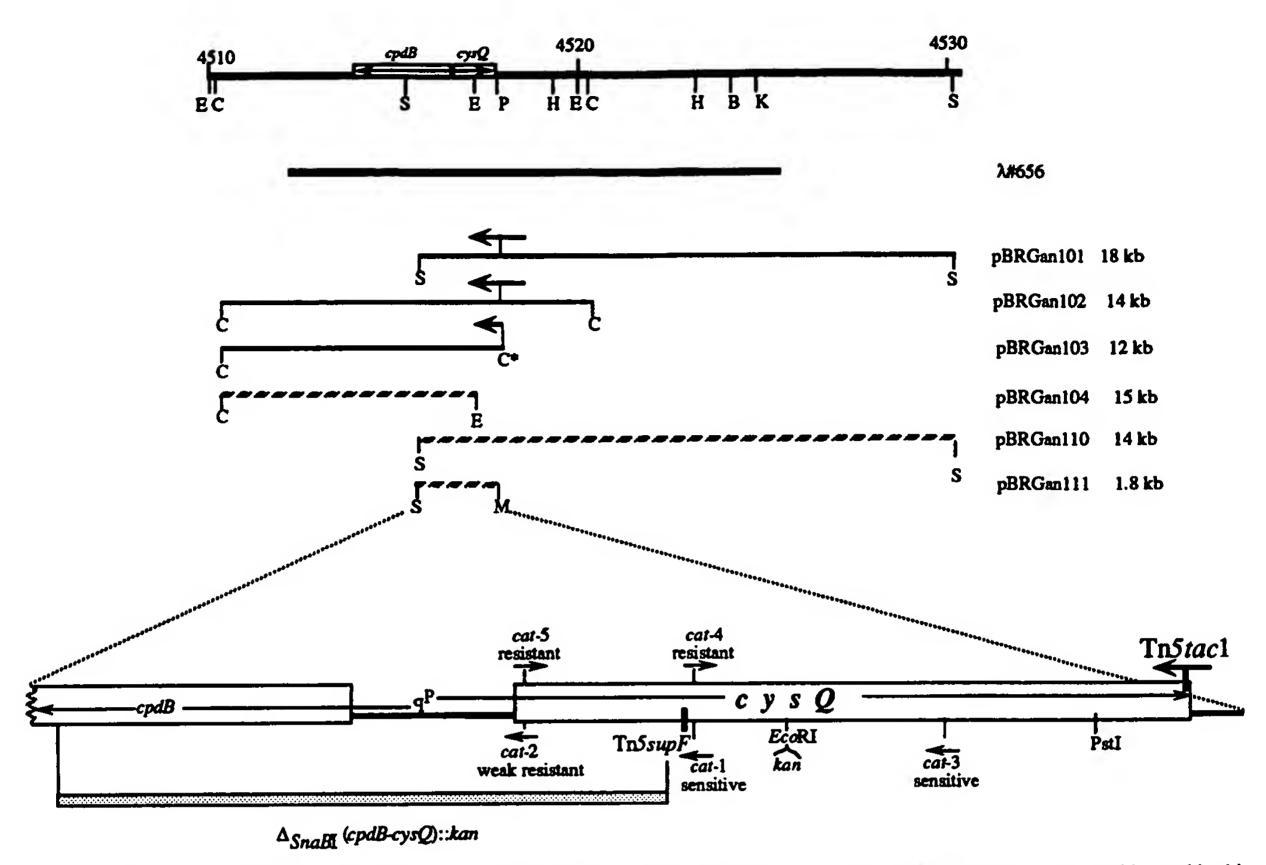


FIG. 2. Physical map of the cysQ region of E. coli. The extents of bacterial DNA cloned in the phage λ656 and in the plasmids used in this work are depicted, as are the positions of insertions and deletions that helped define cysQ. E, EcoRI; C, ClaI; S, SalI; H, HindIII, B, BamHI; K, KpnI. The arrows indicate directions of transcription: heavy arrow, from the tac promoter in Tn5tacI; long arrows, cysQ and cpdB; smaller arrows, orientations of transcription of a cat reporter inserted at Sau3A sites in cysQ.

which contains a chromosomal segment including cpdB and the Tn5tacl insertion. These results implied that the cysteine requirement was due to altered expression of a previously unknown gene next to cpdB. This gene was designated cysQ.

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Direction of cysQ transcription. Insertions of a cat reporter gene were made to determine the orientation of cysQ and thereby to deduce whether auxotrophy resulted from overexpression or underexpression of cysQ after IPTG-induced transcription from the tac promoter in mutant strain DBan41. Five different insertions into Sau3A sites of plasmid pBRGan111 that inactivated Cys+ complementation activity were isolated; restriction mapping showed that each was within about 600 bp of the start of cpdB (Fig. 2). Insertions 4 and 5, oriented away from cpdB (toward Tn5tac1), conferred chloramphenicol resistance (25 µg/ml), whereas insertions 1 and 3, in the opposite orientation, did not. Insertion 2, also in the opposite orientation but closest to cpdB, conferred weak resistance (~10 μg/ml). This was attributed to a second promoter in cysQ that could allow cpdB transcription (see sequence analysis, below). Based on insertions 1, 3, 4, and 5, we inferred that cysQ is transcribed toward Tn5tac1.

Phenotypes conferred by cysQ null alleles. Chromosomal cysQ null mutations were generated and used to assess whether the distinctive leaky auxotrophy and its correction by anaerobic growth were allele or gene specific. (i) A

cysQ::Tn5supF insertion allele was obtained by selecting transposition of Tn5supF to the cysQ⁺ phage λ 656 (29, 43). One of 50 Tn5supF insertions resulted in cysteine auxotrophy when recombined into the E. coli chromosome, and DNA sequencing (see below) showed that Tn5supF was inserted in cysQ. (ii) A cysQ null allele marked with kanamycin resistance was made by insertion of a kan gene at the EcoRI site in pBRGan111 (Fig. 2), recombined from the plasmid into $\lambda656$, and then recombined from the cysQ::kan phage into bacterial chromosomes (28). (iii) A segment containing part of both the cpdB and the cysQ genes was deleted (\Delta SnaBI; Fig. 2) to further test a possible involvement of cpdB in the cysQ mutant phenotype. This deletion was marked by insertion of kan and recombined into $\lambda 656$ and from there into bacterial chromosomes. Finally, E. Barnes kindly provided us with a fourth cysQ null allele, which was generated by Tn10 (Tet) insertion and was originally designated amtA::Tn10 (21, 22).

Each of these four cysQ null alleles resulted in a cysteine requirement that was leaky, corrected by anaerobiosis, and satisfied by sulfite when transduced into several E. coli K-12 strain backgrounds, including DB747 (used to isolate the original Tn5tacl insertion), DB5508, and MG1655. These alleles were much less leaky in two other laboratory strains: TG1 and ET8000. The strain background determines the leakiness of the cysQ mutant phenotype; cysQ derivatives of

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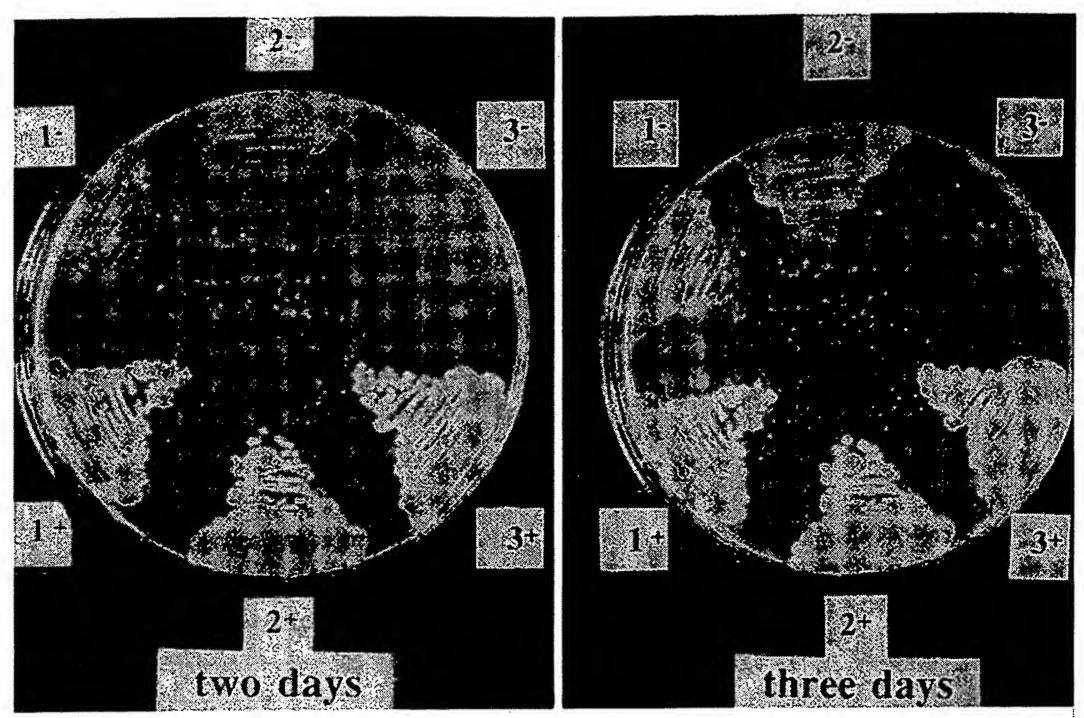


FIG. 3. Growth of cysQ::kan mutants and wild-type parents. Growth was for 2 or 3 days at 37°C on M9 salts glucose medium (no added cysteine or sulfite). (Scattered white dots near the center of plate are a salt precipitate that often form in M9 solid medium.) The numbers 1, 2, and 3 indicate strain backgrounds DB747, MG1655 and ET8000, respectively. –, cysQ::kan; +, Cys⁺.

DB747 and MG1655 showed some growth on cysteine-free medium after 2 days at 37°C and small single colonies after 3 days, whereas the corresponding derivative of ET8000 showed barely perceptible growth only after 3 days (Fig. 3). All cysQ mutant strains grew normally on sulfite or under anaerobic conditions. The Cys^- phenotype caused by the $\Delta(cysQ\ cpdB)$ allele was identical to that caused by simple insertions in cysQ in these strain backgrounds.

The match between the null mutant phenotype and that of the Tn5tacl insertion indicated that IPTG-induced transcription from Tn5tacl shuts off the expression of cysQ quite completely. The equivalence of Cys^- phenotypes of the $\Delta(cysQ\ cpdB)$ and the simple cysQ::kan insertion alleles ruled out a model in which CpdB protein would consume PAPS and in which CysQ protein would regulate this consumption.

DNA sequence of cysQ. A 1-kb segment containing the sites of insertion mutations that defined cysQ was sequenced by using primer binding sites provided by the Tn5tacl and Tn5supF transposons, the kan and cat insertions, and, for one segment, an M13mp18 vector. All portions of this segment were sequenced on both DNA strands. The cysQ DNA sequence corresponded to a 246-codon open reading frame preceded by sequences that match consensus transcription promoters and translation initiation sites (Fig. 4). The first part of this sequence matched that found earlier in the 0.5 kb upstream of cpdB in E. coli; analyses of S. typhimurium indicated a cysQ homolog in the same location in this species (35). The cysQ sequence we determined was identical to that reported for amtA (15).

The DNA sequence confirmed that transcription of cysQ should diverge from that of cpdB, as suggested by cat

reporter insertions. Tn5tac1 was inserted within cysQ, just two codons from its 3' end (a fusion protein with 17 additional amino acids is predicted). Tn5supF was inserted at codon 72 of cysQ. Only 17 bp separate the -35 regions of putative promoters for cysQ and cpdB, and an apparent consensus cyclic AMP receptor protein (CRP) binding site (placed such that CRP binding could stimulate cpdB transcription; see Fig. 4 legend) (35) overlaps the putative cysQ promoter. The low-level chloramphenicol resistance associated with the cat insertion 2 is attributable to an additional promoter within cysQ (nucleotides 72 to 46, underlined in Fig. 4). (This promoter might also explain the relative weakness of the CRP-cyclic AMP dependence of cpdB expression [35].) Binding sites for the CysB positive regulatory protein, found in the promoter regions of most cysteine biosynthetic genes (18), did not seem to be present in the cysQ promoter region. A search of the PROSITE data base of protein motifs (2) did not reveal significant matches to CysQ. However, we recently found strong amino acid sequence level homologies between cysQ and suhB of E. coli (Fig. 5), mutations in which suppress certain rpoH missense alleles (apparently by elevating the levels of heat shock sigma subunit of RNA polymerase that it encodes [56]) and also between cysQ and genes for several eukaryotic proteins, including inositol monophosphatase (13, 39).

Does cysQ participate in ammonium uptake? While preparing this manuscript, we learned that cysQ corresponds to the gene called amtA (21, 22), a designation based on the finding of a Tn10 insertion mutation (amtA::Tn10) that blocked growth on minimal (cysteine-free) medium containing very low levels of ammonium (≤ 0.1 mM, rather than the ≥ 10 mM used in most media). It was proposed that amtA is needed

SD(cpd) AAT<u>CAT</u>CAGGGACA<u>TCCTT</u>TTATCATCGGGAATACGAAAGAAAAGGGAGAATAAACGTCT -35(cpd) -10(cpd) TACTTATAGAACAGTGAAGAATGCCACAATTTTACGCTTTGAAAATGATGACACTATCAC -10(cys)-35(cys) AGTTGGCGCATTCATTAACGATAGGGTATAAGTAAAACAATAAGTTAACACCGCTCACAG CAT 25 AGACGACGTGGAGAAATGTTAGATCAAGTATCCCAGCTTGCACGGAATGCAGGCGATGCC MetLeuAspGlnValCysGlnLeuAlaArgAsnAlaGlyAspAla 46 <u>ATTATGCAGGTCTACGACGGACGAAACCGATGGACGTCGTCAGCAAAGCGGACAATTCT</u> IleMetGlnValTyrAspGlyThrLysProMetAspValValSerLysAlaAspAsnSer ProValThrAlaAlaAspIleAlaAlaHisThrValIleMetAspGlyLeuArqThrLeu **CAT 1.4** (1)Tn5supF(O)166 ACACCGGATGTTCCGGTCCTTTCTGAAGAAGATCCTCCCGGTTGGGAAGTCCGTCAGCAC ThrProAspValProValLeuSerGluGluAspProProGlyTrpGluValArgGlnHis 226 TGGCAGCGTTACTGGCTGGTAGACCCGCTGGATGGTACTAAAGAGTTTATTAAACGTAAT TrpGlnArgTyrTrpLeuValAspProLeuAspGlyThrLysGluPheIleLysArgAsn **EcoRI** 286 GGCGAATTCACCGTTAACATTGCGCTCATTGACCATGGCAAACCGATTTTAGGCGTGGTG GlyGluPheThrValAsnIleAlaLeuIleAspHisGlyLysProIleLeuGlyValVal 346 TATGCGCCGGTAATGAACGTAATGTACAGCGCGGCAGAAGGCAAAGCGTGGAAAGAAGAG TyrAlaProValMetAsnValMetTyrSerAlaAlaGluGlyLysAlaTrpLysGluGlu CysGlyValArgLysGluIleGlnValArgAspAlaArgProProLeuValValIleSer 466 CGTTCCCATGCGGATGCGGAGCTGAAAGAGTATCTGCAACAGCTTGGCGAACATCAGACC ArgSerHisAlaAspAlaGluLeuLysGluTyrLeuGlnGlnLeuGlyGluHisGlnThr 526 ACGTCCATCGCTCTTCGCTGAAATTCTGCCTGGTGGCGGAAGGACAGGCGCACGTGTAC ThrSerIleGlySerSerLeuLysPheCysLeuValAlaGluGlyGlnAlaHisValTyr 586 CCGCGCTTCGGACCAACGAATATTTCGGACACCGCCGCTGGACATGCTGTAGCTGCAGCT ProArgPheGlyProThrAsnIleTrpAspThrAlaAlaGlyHisAlaValAlaAlaAla 646 GCCGGAGCGCACGTTCACGACTGGCAGGGTAAACCGCTGGATTACACTCCGCGTGAGTCG AlaGlyAlaHisValHisAspTrpGlnGlyLysProLeuAspTyrThrProArgGluSer 741 (O)Tn5tac1(I) 706 TTCCTGAATCCGGGGTTCAGAGTGTCTATTTACTAAATTCAGATGGCAGAAACAGTGTAT PheLeuAsnProGlyPheArgValSerIleTyrEnd

TATTTAAAGTGCAAAAATTCAATTGCTAATAAGTTACA

FIG. 4. DNA sequence of the cysQ gene. Sites of insertion and orientations (in cases of transposons) are indicated, as are putative promoters for cysQ and cpdB and the consensus CRP binding site. The CRP binding site identified in ref. 35 extends from positions -72 to -96, relative to the start of cysQ translation (beginning at TT in the -35 region of the cysQ promoter). The DNA sequencing protocols and sequences of the oligonucleotide primers used are given in Materials and Methods.

for active ammonium uptake (21). Our reconstruction experiments showed, however, that even $cysQ^+$ ($amtA^+$) bacteria grew very poorly on low-ammonium medium (Fig. 6). Strains with the amtA::Tn10 or cysQ::kan insertion mutations failed to form colonies on this medium, as reported earlier (21). The mutants did grow, however, when this medium was supplemented with cysteine (Fig. 6) or sulfite, in which case these mutant strains were indistinguishable from their Cys^+ parents in colony size. Since neither sulfite nor cysteine can be used as an ammonium source by $E.\ coli$ (53), these results are not consistent with the interpretation (21) that the AmtA (CysQ) protein is needed for ammonium uptake.

Possible roles for cysQ. We tested whether cysQ might be needed in sulfate uptake or activation. No effect of cysQ null mutations was found on the rapid uptake of labelled sulfate from the medium in the background of strain TG1 or DB5463 (nonleaky and leaky cysteine requirements, respectively). In contrast, a cysDN (ATP sulfurylase) deletion strain was severely deficient in sulfate uptake, as expected (14) (data not shown). No significant differences between cysQ mutant and parental strains were detected in levels of ATP sulfurylase, which catalyzes synthesis of APS, the first activated sulfur intermediate. cysQ mutations also had no effect on the level of the activator of ATP sulfurylase (data not shown).

Reversion of cysQ null mutants. Cys+ revertants of

MLDQVCQLARNAGDAIMQVYDGTKPMDVVSKADNSPVTAADIAAHTVIMDG	(1-51)	CysQ
ML AR AG I Y+ ++ K+ N VT D AA VI+D MHPMLNIAVRAARKAGNLIAKNYETPDAVEASQKGSNDFVTNVDKAAEAVIIDT	(1-54)	SuhB
LRTLTPDVPVLSEEDPPGWEVRQHWQRYWLVDPLDGTKEFIKRNGEFTVNIALI +R P +++EE E W++DPLDGT FIKR F V IA+	(52-105)	CysQ
IRKSYPQHTIITEE-SGELEGTDQ-DVQWVIDPLDGTTNFIKRLPHFAVSIAVR	(55–106)	SuhB
DHGKPILGVVYAPVMNVMYSAAEGKAWKEECGVRK-QIQVRDARPPLVVISRSH +G+ ++VVY P+ N +++A G + G R RD ++	(106-157)	CysQ
IKGRTEVAVVYDPMRNELFTATRGQG-AQLNGYRLLGSTARDLDGTILATGFPF	(107-159)	SuhB
ADAEL KEYLOOLGEHOTTSIGSS-LKFCLVAEGOAHVYPRFGPTNIWD Y++ +G GS L VA G + G WD	(158-205)	CysQ
KAKQYATTYINIVGKLFNECADFRRTGSAALDLAYVAAGRVDGFFEIG-LRPWD	(160-212)	SuhB
TAAGHAVAAAAGAHVHDWQGKPLDYTPRESFLNPGFRVSIY AAG + AG+ V D G Y + RV	(206-246)	CysQ
FAAGELLVREAGGIVSDFTGGH-NYMLTGNIVAGNPRVVKAMLANMRDELSDALKR	(213-267)	SuhB

FIG. 5. Alignment of amino acid sequences of inferred protein products of cysQ and suhB (adapted from data in reference 39). Identities are indicated by placements of conserved amino acids in the middle line; conservative substitutions (+) are indicated. Overlined segments indicate regions with high sequence similarity to inositol monophosphatase (13). The CysQ (SuhB) initial amino acid alignment score of 229, calculated by using the FASTA program (42), was 28 standard deviation units above the mean initial score of 24.6 for comparisons of CysQ to the other sequences in the PIR (release 28) protein data base. In a test using the Dayhoff Relate program, there were four segments of 25 amino acids in length that were more than eight standard deviations above the mean. This indicates strong homology: the probability of getting a single segment with such a deviation from a random sequence by chance alone is less than 10⁻¹⁵ (42).

cysQ::kan and cysQ::Tn10 mutants were obtained at frequencies of about 10⁻⁶ with derivatives of TG1 and ET8000 (nonleaky cysQ mutant phenotype) and $>10^{-5}$ with derivatives of DB747 and MG1655 (leaky cysQ mutant phenotype); these differences in recovery probably reflect the greater leakiness of cysQ mutations in the DB747 and MG1655 backgrounds. The revertants were heterogeneous in colony size on cysteine-free medium but grew as well as their cysQ+ ancestors on cysteine-containing medium and retained the Kan' or Tet' traits of their Cys parents. The parental cysQ mutant alleles were recovered from several revertants by transduction and selection for the appropriate resistance trait (Kan' or Tet'). Two spontaneous reversion mutations that allowed relatively good growth on cysteine-free medium were mapped in Hfr × F crosses and then by transduction with several candidate λ phage clones (marked by insertion of a Tn5cam transposon [44, 50]). These reversion mutations were found to be in the segment carried by $\lambda 419$, a phage clone that also carries the cysPTWA (sulfate binding and uptake) operon. The two revertants tested were found to be defective in sulfate uptake, unlike their cysteine-requiring parents. Partial diploids generated by lysogenizing revertants with $\lambda 419$::Tn5cam and a λ^+ helper required cysteine, indicating that the reversion mutation is recessive and thus probably due to loss of function. This Cys phenotype was unstable, however, because of frequent homogenotization for the parental (nonrevertant) allele.

DISCUSSION

The initial steps in the sulfate assimilation branch of the cysteine pathway entail sulfate uptake, its activation via formation of APS and conversion to PAPS, and then its reduction to sulfite (Fig. 1). The mutational and sequence analyses presented here identified a previously uncharacterized gene, cysQ, whose product is needed for proper metabolic functioning of this part of the pathway. We propose below that CysQ acts on PAPS. The cysQ gene was mapped

to a locus at ~ 96 min in the $E.\ coli$ chromosome, which is far from other cys genes. The cysQ promoter region overlapped a CAP binding site that is implicated in the control of expression of the adjacent gene, cpdB (35), and it did not contain a good match to the consensus binding site for the CysB regulatory protein (18). Hence, the expression of cysQ may be controlled differently from that of most other cys genes. The auxotrophy resulting from mutations in cysQ was leaky in some strain backgrounds and was compensated by mutations in other genes; cysQ mutants were prototrophic during anaerobic growth.

A precedent for cysteine biosynthetic genes that are not needed during anaerobic growth is provided by cysI and cysJ in S. typhimurium (3). This case reflects the presence of additional anaerobic sulfite reduction (asr) genes. E. coli lacks such asr genes (20), however, and our studies indicate that cysQ acts before, not after, sulfite formation (Fig. 1). cysB is also only needed during aerobic growth (3), suggesting that a separate transcriptional activator may be operating anaerobically. cysQ does not seem to be a transcriptional activator of cys genes, since it does not significantly affect the rate of sulfate uptake or the level of ATP sulfurylase or its protein activator.

cysQ is identical to amtA, a gene which had been thought to participate in ammonium transport (21). That interpretation was based on a failure of mutant strains to grow on low-ammonium cysteine-free medium or to take up methylammonium at a high rate when they were grown with arginine in place of ammonium (21). We found that the growth defect of cysQ (amtA) mutants was compensated by sulfite or cysteine (Fig. 6), neither of which serves as a nitrogen source (53). The initial failure to recognize the cysteine requirement of the amtA mutant (21) may have been due to leakiness on normal minimal medium (Fig. 3) or inadvertent selection of a (partially) compensating suppressor mutation. The inability of cysQ (amtA) mutants to grow on low-ammonium medium probably results from the combined effects of partial starvation for ammonium (because of

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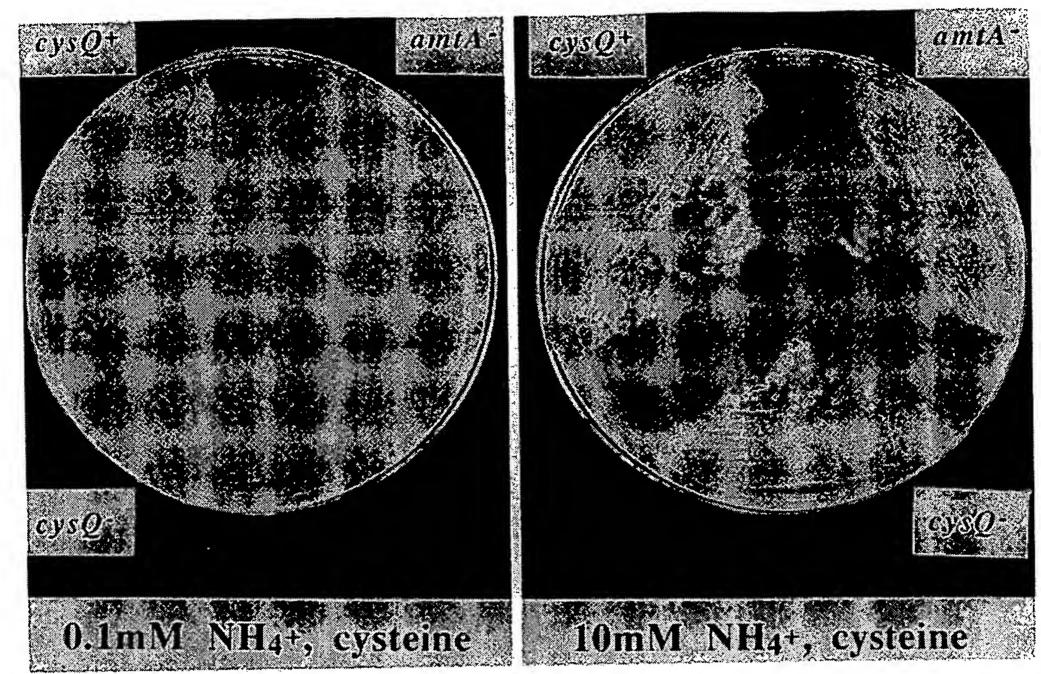


FIG. 6. Comparison of wild-type and cysQ and amtA mutant bacterial strains on modified M9 minimal medium and either low (0.1 mM) or normal (10 mM) concentrations of ammonium acetate. Strains: cysQ⁺, ET8000; cysQ mutant, ET8000 cysQ::kan (DB6908); amtA mutant, ET8000 amtA::Tn10 (AJ2653) grown in medium containing 0.2 mM cysteine. The plate with 10 mM ammonium was incubated for 1 day and the plate with 0.1 mM ammonium was incubated for 2 days at 37°C before being photographed. Identical growth patterns were obtained with sulfite in place of cysteine, but only Cys⁺ parental strains grew on minimal low-ammonium medium lacking sulfite or cysteine. Equivalent weak growth of mutant and Cys⁺ sibling strains on the low-ammonium medium was also observed with all other lineages tested (DB747 and its cysQ derivative DB5656, MG1655 and its cysQ derivative DB6316, TG1 and its cysQ derivative DB6913). Growth was weaker on medium containing 0.025 mM rather than 0.1 mM ammonium acetate, as expected, since ammonium was limiting. In the cases of Cys⁺ strains, this slow growth was not stimulated by adding 10 or 20 mM cysteine (or sulfite or thiosulfate), whereas growth was stimulated by adding 20 mM glutamate or arginine (which serve as ammonium sources [53]).

the medium) and for cysteine (because of a mutation). Although the inefficient induced methylammonium uptake by amtA cells was also interpreted to reflect a specific uptake defect, the reported data (21) indicate that the basal level of uptake was not affected by the amtA mutation. Earlier work had shown that induction by growth in arginine reflects the slow release of ammonium from this source, relative to the rate of ammonium consumption (45, 53). Because the poor growth of cysQ (amtA) mutants on cysteine-free medium should allow the arginine-derived ammonium to accumulate to repressing levels, we do not find it necessary to postulate a role for CysQ (AmtA) in ammonium or methylammonium uptake.

How does CysQ act in the synthesis of sulfite and cysteine? Several possible roles have been eliminated by our results to date: (i) sulfate uptake, (ii) stabilization of ATP sulfurylase, (iii) synthesis or stabilization of the ATP sulfurylase activator, and (iv) modulation of CpdB. In addition, the cysQ sequence does not match that of ppa, the gene for pyrophosphatase (30), an enzyme probably needed for efficient APS synthesis (Fig. 1). The leakiness of cysQ-null alleles in many strain backgrounds might reflect (i) a second gene with a functionally related role; (ii) an intrinsic activity of gene(s) that can mutate to give a Cys⁺ revertant phenotype; or, if cysQ is regulatory, (iii) strain backgrounddependent differences in the quantitative effects of CysQ on the gene, protein, or metabolite that is the target of its control.

Studies of Cys⁺ revertants are providing insights into how CysO may act. Several spontaneous reversion mutations were mapped to a region that includes cysTWA permease genes, were recessive to the wild-type (nonrevertant) alleles, and were defective in sulfate uptake. These results suggested that reversion results from loss of function, not from an unusual expression of a silent or cryptic suppressor gene. Accordingly, we have begun to isolate transposon insertions that restore prototrophy to cysQ mutants in a nonleaky background (50). One insertion that resulted in very small colonies on cysteine-free medium was in cysA, which encodes a subunit of sulfate permease (49). Transduction of this insertion into a $cysQ^+$ strain resulted in the same small-colony phenotype, indicating that the phenotype reflected loss of cysA function, not poor suppression of the cysQ mutation. A second insertion, which resulted in colonies of nearly normal size, was in cysP, a gene whose product contributes to efficient sulfate and thiosulfate binding (19). In interpreting these reversion data we draw on early findings that mutations in cysH or in trxA plus grx cause poor growth, apparently because accumulated PAPS or a derivative of it is toxic, and that these mutations can be compensated by mutations inactivating sulfate permease (16, 45a). Although the role of cysP in the cys pathway is not understood, the ability of permease mutations to compensate for the defect in cysQ suggests that CysQ also acts on PAPS. Perhaps CysQ participates with the CysH sulfotransferase to generate sulfite. Alternatively, perhaps CysQ sequesters or consumes excess PAPS or a toxic derivative of it. On this latter view, cysteine might be needed for growth of cysQ mutants only to allow repression of cys gene expression and thereby decrease PAPS synthesis, rather than to compensate for a missing biosynthetic enzyme. CysQ exhibits striking amino acid sequence homology to mammalian inositol monophosphatase as well as to the product of the suhB gene of E. coli (Fig. 6) (39). The homology between CysQ and inositol monophosphatase, in particular, encourages models in which CysQ acts on a phosphorylated metabolite such as PAPS, possibly ensuring that it plays its essential biosynthetic role without toxicity to the cell.

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Identification of a Third Sulfate Activation System in *Sinorhizobium* sp. Strain BR816: the CysDN Sulfate Activation Complex

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Sinorhizobium sp. strain BR816 possesses two nodPQ copies, providing activated sulfate (3'-phosphoadenosine-5'-phosphosulfate [PAPS]) needed for the biosynthesis of sulfated Nod factors. It was previously shown that the Nod factors synthesized by a nodPQ double mutant are not structurally different from those of the wild-type strain. In this study, we describe the characterization of a third sulfate activation locus. Two open reading frames were fully characterized and displayed the highest similarity with the Sinorhizobium meliloti housekeeping ATP sulfurylase subunits, encoded by the cysDN genes. The growth characteristics as well as the levels of Nod factor sulfation of a cysD mutant (FAJ1600) and a nodP1 nodQ2 cysD triple mutant (FAJ1604) were determined. FAJ1600 shows a prolonged lag phase only with inorganic sulfate as the sole sulfur source, compared to the wild-type parent. On the other hand, FAJ1604 requires cysteine for growth and produces sulfate-free Nod factors. Apigenin-induced nod gene expression for Nod factor synthesis does not influence the growth characteristics of any of the strains studied in the presence of different sulfur sources. In this way, it could be demonstrated that the "household" CysDN sulfate activation complex of Sinorhizobium sp. strain BR816 can additionally ensure Nod factor sulfation, whereas the symbiotic PAPS pool, generated by the nodPQ sulfate activation loci, can be engaged for sulfation of amino acids. Finally, our results show that rhizobial growth defects are likely the reason for a decreased nitrogen fixation capacity of bean plants inoculated with cysD mutant strains, which can be restored by adding methionine to the plant nutrient solution.

Sulfur is a macronutrient that is required by all organisms. It forms constituents of proteins, lipids, carbohydrates, electron carriers, and numerous cellular metabolites. Sulfate is the most abundant source of utilizable sulfur in the aerobic biosphere. The sulfate assimilation complex, required for the formation of the sulfur-containing amino acid cysteine, has been the subject of intensive study in *Escherichia coli* (21). Cysteine is the central precursor of all organic molecules containing reduced sulfur, ranging from the amino acid methionine to peptides, proteins, vitamins, cofactors such as S-adenosylmethionine, and hormones.

Like all inorganic nutrients, sulfate is transported into cells by highly specific membrane transport systems (18). Sulfate assimilation requires its prior activation to adenylate compounds via a pathway that seems to be similar in all organisms. The activation is achieved by the ATP sulfurylase-catalyzed reaction of sulfate with ATP to give adenosine 5'-phosphosulfate (APS), coupled with GTP hydrolysis. Subsequently, APS is phosphorylated by an APS kinase to produce 3'-phosphoadenosine-5'-phosphosulfate (PAPS). In E. coli, ATP sulfurylase is encoded by cysD and cysN, whereas the APS kinase is encoded by cysC (27, 28). PAPS is then enzymatically reduced by the cysH-encoded PAPS reductase (also known as PAPS sulfotransferase) to sulfite, which enters the cysteine biosynthetic pathway.

PAPS also serves directly as a sulfate donor for the forma-

tion of sulfated compounds. For example, Rhizobium-legume symbiotic interactions are mediated by a host-specific bacterial signaling molecule (the Nod factor), which can be sulfated. In general, rhizobial species that produce sulfated Nod factors possess at least two sulfate activation systems (6, 12, 24, 25, 40). The three genes that are indispensable for Nod factor sulfation, nodP, nodQ, and nodH, were first isolated from Sinorhizobium meliloti. Together, nodP and nodQ encode both ATP sulfurylase and APS kinase activities (45, 47), whereas the nodH gene product, a sulfotransferase, directly transfers the activated sulfate moiety to the Nod factor backbone (8, 44). NodP is homologous to E. coli CysD, while the amino- and carboxy-terminal domains of NodQ are homologous to E. coli CysN and CysC, respectively. In a recent study, it was reported that the specificity of phytopathogen-host interactions also can be controlled by a sulfated avirulence effector molecule, which is yet to be identified (48). The rice pathogen Xanthomonas oryzae pv. oryzae RaxP and RaxQ proteins are responsible for the synthesis of an activated form of sulfate and are similar to the NodP and NodQ host specificity proteins of the bacterial symbiont S. meliloti.

In S. meliloti, two copies of the nodPQ operon are present. Both copies are involved in Nod factor sulfation but are not necessary for cysteine biosynthesis. Recently, in S. meliloti and in Rhizobium tropici CFN299, homologues of the cysDN (ATP sulfurylase) and cysH (APS reductase) genes were isolated, but no homologue of the E. coli cysC gene (APS kinase) could be identified (1, 23). Consequently, it was demonstrated that in S. meliloti, APS rather than PAPS is reduced for sulfite production during cysteine biosynthesis (1). Other members of the

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TABLE 1. Bacterial strains and plasmids

Strain or plasmid	Relevant characteristics	Reference or source	
Sinorhizobium sp. strains			
BR816	Broad-host-range Sinorhizobium strain isolated from Leucaena leucocephala	16	
FAJ1600	cysD mutant of BR816; Tc ^r	This study	
FAJ1604	nodP1 nodQ2 cysD triple mutant of BR816; Km ^r Sp ^r Tc ^r	This study	
CFNE205	nodP1 mutant of BR816; Km ^r	25	
CFNE206	nodQ2 deletion mutant of BR816; Sp ^r	25	
CFNE207	nodP1 nodP2 double mutant of BR816; Km ^r Sp ^r	25	
CFNE208	nodPl nodQ2 double mutant of BR816; Km ^r Sp ^r	T. Laeremans, unpublished result	
Plasmids			
pBRE4.8	pUC19 carrying the BR816 cysDN genes; Apr	This study	
pJQ200uc1	B. subtilis sacB-containing suicide vector; Gm ^r	39	
pHP45Ω-Tc	Vector containing Tc ^r cassette	38	
pUC18/19	Cloning vector; Apr	33	

Rhizobiaceae, differing in their ability to incorporate sulfate in either a Nod factor or lipopolysaccharide, also preferentially reduce APS instead of PAPS for cysteine biosynthesis. This implies that APS reduction is not necessarily correlated with the presence of PAPS-dependent sulfurylation reactions for symbiosis, which is the case when functional nodPQ genes are present (1). Recently, Kopriva et al. (20) have described a phylogenetic classification of APS and PAPS reductase amino acid sequences (both annotated as CysH) from different organisms. The resulting sequence-based prediction of the substrate specificities of these enzymes was confirmed by Williams et al. (58), using genetic complementation experiments.

Sinorhizobium sp. strain BR816 (formerly Rhizobium sp. strain BR816) synthesizes Nod factors that are fully sulfated at the reducing terminal residue (50), as is the case for the narrow-host-range S. meliloti (26). The sulfate decoration on the Nod factors secreted by S. meliloti is essential for nodulation of alfalfa (40). Except for S. meliloti, it is still unclear whether rhizobia producing sulfated Nod factors use only the nodPQdependent PAPS pool as a source of activated sulfate for Nod factor sulfation, the housekeeping PAPS pool, or both (25). Previously, Laeremans et al. (25) demonstrated that Sinorhizobium sp. strain BR816 possesses two nodPQ copies. Although both copies are functional, as demonstrated by genetic complementation of an R. tropici nodP mutant, the double mutants did not show any detectable changes in the amount of sulfated Nod factors produced by this strain (25). It was suggested that in Sinorhizobium sp. strain BR816, in contrast to S. meliloti, a housekeeping locus as a third PAPS-producing locus could be involved in the sulfation of the Nod factors.

We have isolated the cysDN homologues of Sinorhizobium sp. strain BR816 and studied the role of this third PAPS-producing locus in relation to Nod factor synthesis. In addition, we were interested to know how the various forms of activated sulfate may be partitioned into the pathways for amino acid biosynthesis and sulfation or methylation of Nod factors and other compounds important during symbiosis. Furthermore, based on the analysis of the phylogenetic relationship among rhizobial ATP sulfurylases, we speculate on the possible origin and functionality of genes for sulfate activation.

MATERIALS AND METHODS

Bacterial strains and growth conditions. The bacterial strains and plasmids used in this study are listed in Table 1. E. coli strains were maintained on Luria-Bertani agar at 37°C and grown in Luria-Bertani broth (32). Rhizobial strains were maintained on yeast extract-mannitol medium (55) or on tryptone-yeast medium with added CaCl₂ (3) at 30°C. Antibiotics were added to the medium as needed at the following concentrations (micrograms per milliliter): ampicillin. 100; spectinomycin, 50; kanamycin, 50; and nalidixic acid, 31. Tetracycline was added to a final concentration of 1 µg/ml (for Sinorhizobium sp. strain BR816) or 10 µg/ml (for E. coli). Triparental conjugations and site-directed mutagenesis were done as previously described (31).

Nucleic acid manipulations and analysis. Isolation and cloning of plasmid DNA was performed as described previously (2, 42). Total genomic DNA of Sinorhizobium sp. strain BR816 was isolated by using a genomic DNA isolation kit (Gentra Systems) according to the manufacturer's instructions. DNA fragments were recovered from agarose gels by using the Nucleotrap kit (Macherey-Nagel). Southern blotting and hybridizations were carried out as previously described (25). Sequencing of DNA fragments cloned in the pUC18-pUC19 vectors was performed on an automated ALF sequencer with fluorescein-labeled universal and synthetic oligonucleotide primers (Amersham Pharmacia Biotech, Uppsala, Sweden). Database searches for similarity were performed with the BLAST software (National Center for Biotechnology Information, National Institutes of Health).

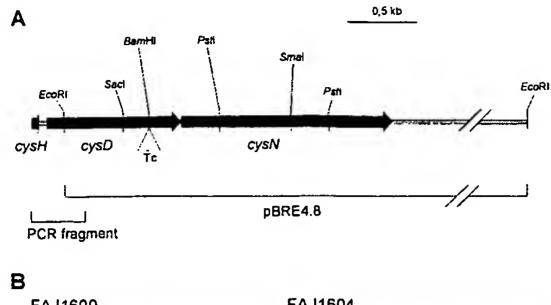
PCR was performed with Taq DNA polymerase (Bochringer, Mannheim, Germany) according to the manufacturer's protocol. For sequencing, the high-fidelity Platinum Pfx DNA polymerase (GIBCO-BRL, Life Technologies) was used according to the manufacturer's protocol.

To construct a genomic minilibrary, total genomic DNA from Sinorhizobium sp. strain BR816 was digested with EcoRI. DNA fragments ranging between 4 and 6 kb were recovered and ligated into the pUC19 cloning vector. Eight hundred Apt white colonies were picked up. Plasmid DNA was purified from 15 pools consisting of approximately 50 colonies, and efficient insertion of fragments of the desired size was confirmed. A 450-bp PCR fragment containing an internal part of cysD was used as a probe to screen the library.

Phylogenetic analysis of CysD homologues. The amino acid sequences of 19 CysD-like proteins, truncated to the same size as the shortest sequence (position 3 to 299 from the S. meliloti NodP1 sequence [gi14523565]) were aligned by using the ClustalW program (http://searchlauncher.bcm.tmc.edu/multi-align/multi-align.html). The construction of neighbor-joining trees (41) and bootstrap analysis of 1,000 resamples were performed by using the Treecon for Windows (1.3b) software package (53). In estimating evolutionary distances between amino acid sequences, we used the Poisson correction. Insertions and deletions were not taken into account. For constructing trees by the parsimony method, the PROTPARS program in the PHYLIP package was used (10). Again, bootstrap analysis of 1,000 resamples was performed.

Growth tests. Growth tests of Sinorhizobium sp. strain BR816 in sulfate-free liquid medium were carried out in acid minimal salts (AMS) medium (36) containing 1 mM CaCl₂ with sulfate salts replaced by equimolar amounts of alternative salts (MgCl₂, ZnCl₂, MnCl₂, and CuCl₂). Ammonium chloride (10)

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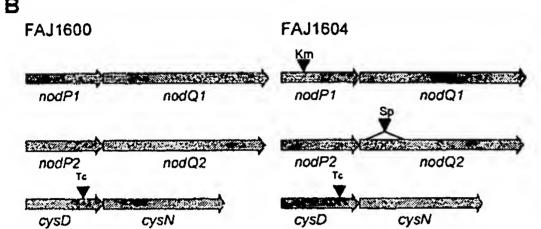


FIG. 1. (A) Physical and genetic maps of the BR816 cysDN region. The triangle indicates the position of the inserted Tc^r cassette in the mutants FAJ1600 and FAJ1604. (B) Schematic view of the constructed Sinorhizobium sp. strain BR816 mutants with mutations in the nodPQ genes and cysDN genes (see Table 1). Triangles indicate inserted antibiotic resistance cassettes.

mM) and mannitol (10 mM) were used as nitrogen and carbon sources, respectively. Sulfur compounds (sodium sulfate, sodium sulfite, 1.-cysteine, and 1.-methionine) were filter sterilized and added to the autoclaved medium at a concentration of 25 μM. When appropriate, cell cultures were induced with 500 nM apigenin. Cells of the strains tested were grown overnight in tryptone-yeast medium, washed twice in sulfate-free AMS medium, brought to an optical density of 0.4 (measured at 600 nm with a Perkin-Elmer lambda 2 spectrometer), and diluted 6,000-fold in sulfate-free AMS medium with the appropriate concentrations of filter-sterilized antibiotics, apigenin, and sulfur compounds. Bacteria were grown in microtiter plates (final volume, 300 μl) over a 4-day period, and cell growth was monitored automatically by measuring the optical density at 600 nm in BioscreenC (Labsystems) every 30 min. For each time point, the average optical density was calculated from five independent measurements.

Insertion mutagenesis. A Sinorhizobium sp. strain BR816 cysD single mutant and nodP1 nodQ2 cysD triple mutant were constructed as follows. To obtain the cysD single mutant, the 1.6-kb Sma1 fragment of pBRE4.8 was ligated into the Sma1 site of pJQ200uc1. This vector allows positive selection of double homologous recombinants on sucrose (10%)-containing medium due to the presence of the Bacillus subtilis sacB gene. The resulting plasmid was digested with BamHI and then blunt-end ligated to the Sma1 fragment containing the Ω -Tc^r cassette from pHP45 Ω -Tc. This plasmid was conjugated to Sinorhizobium sp. strain BR816. Correct insertion of the Tc^r interposon was verified by Southern hybridization with the cysD gene and the Tc^r cassette as probes. In this way, the same construct was introduced in CFNE205 (nodP1), CFNE206 (nodQ2), CFNE207 (nodP1 nodP2), and CFNE208 (nodP1 nodQ2) (Table 1; Fig. 1). A cysD single mutant (FAJ1600) and a nodP1 nodQ2 cysD triple mutant (FAJ1604) were obtained and retained for further analysis.

Radioactive labeling of Nod metabolites and thin-layer chromatography (TLC) analysis. Nod factors were labeled by using the isotopes [14C]acetate and [35S]sulfate according to a slightly modified version of the protocol of Mergaert et al. (30), as previously described (25). For this experiment, Nod factors were purified from cells grown in sulfate-free AMS minimal medium supplemented with L-cysteine, as described for the growth tests.

Plant nodulation assay. Seeds of *Phaseolus vulgaris* ev. BAT477 were surface sterilized and germinated as described previously (56). Bean seedlings were planted in 250-ml flasks containing a nitrogen-free Snoeck medium agar slant (C. Snoeck, J. Vanderleyden, and E. Schrevens, submitted for publication) with KH₂PO₄ (7.49 mM), K₂SO₄ (0.43 mM), CaCl₂ (2.65 mM), MgCl₂ (1.75 mM), MgSO₄ (1.2 μM), FeNaEDTA (50.8 μM), MnSO₄ (35.2 μM), CuSO₄ (0.5 μM), ZnSO₄ (1.5 μM). H₃BO₃ (25 μM), and (NH₄)₆Mo₇O₂₄ (0.07 μM), with sulfate as the sole sulfur source unless otherwise stated. The seedlings were inoculated

with approximately 10⁶ bacteria per plant, from a diluted overnight culture that was washed twice with sulfate-free AMS medium. The plants were maintained in a growth chamber at 26°C (day) and 22°C (night) with a 12-h photoperiod. Plants were harvested after 3 weeks. Uninoculated control plants did not show any nodules or nodule-like structures. Ten plants per strain were tested in each experiment. Nitrogenase activity was determined by measuring the acetylene reduction activity of nodulated roots in closed vessels with a Hewlett-Packard 5890A gas chromatograph equipped with a PLOT fused silica column, with propane as an internal standard.

Data analysis. In all experiments, a randomized block design was used with 10 replicate blocks. Nodule number, nodule dry weight, and acetylene reduction activity were analyzed with the means and general linear model procedure (SAS Institute, Cary, N.C.). Comparison among the mean values obtained for each strain was made by Tukey's multiple-range test with a 95% confidence limit.

Nucleotide sequence accession number. Nucleotide sequence data were deposited in the GenBank database under accession number AJ505754.

RESULTS

Cloning and sequencing of a third PAPS-producing locus in Sinorhizobium sp. strain BR816. Previous work provided evidence for the presence of a putative third PAPS-producing locus in Sinorhizobium sp. strain BR816 on an approximately 4.8-kb EcoRI genomic DNA fragment (25). In order to clone this third copy of sulfate activation genes, a genomic minilibrary was constructed (see Materials and Methods), and a single positive clone, pBRE4.8, was obtained. Since the inserted genomic DNA region corresponding to the cysD gene was incomplete, the missing part of cysD was obtained by PCR with primers that were designed based on existing knowledge of the genomic organizations and DNA sequences of sulfate assimilation genes in other Rhizobium spp. (1, 23).

A physical map of the 4.8-kb EcoRI fragment and the upstream 442-bp PCR fragment was established (Fig. 1A), and the nucleotide sequence was determined. Similarity with an ATP sulfurylase encoded by the cysD and cysN genes of S. meliloti, R. tropici CFN299, and E. coli was found. Partial sequence similarity upstream of the cysD gene revealed the presence of a cysH homologue, encoding an APS or PAPS reductase, whereas no cysC homologue was found in the sequenced fragment. The same organization is found in S. meliloti and R. tropici (1, 23). It is likely that all three open reading frames are in a single operon, since no promoter consensus sequences or transcription termination signals were found in the intergenic cysH-cysD sequence of BR816. A similar situation was observed in S. meliloti, where two transcriptional start sites were identified, both upstream of the cysH homologue (1). In contrast, in E. coli, cysH does not form an operon with cysDNC (21). The nodP and nodQ homologues have a lower percent G+C content than the cysD and cysN homologues (data not shown), as observed for the S. meliloti genome (13).

The Sinorhizobium sp. strain BR816 cysD and cysN genes encode proteins of 317 and 498 amino acids, respectively. Strong conservation of amino acid residues was found with the respective CysD and CysN proteins of S. meliloti (96 and 91% identity, respectively), R. tropici (89 and 82% identity), and E. coli (68 and 52% identity). CysN contains the characteristic GTP-binding motif (GxxxxGK, DxxG, and NKxD) (7) and also an ITI motif, which is conserved among elongation factors (19). In comparison to the NodQ peptides, the deduced amino acid sequence of cysN lacks the carboxy-terminal part that corresponds to E. coli CysC. Therefore, no ATP-binding or PAPS-binding motifs were found. Similar observations were

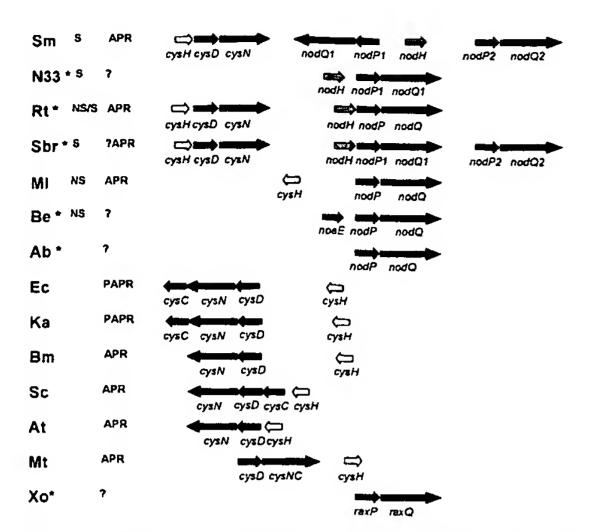


FIG. 2. Schematic representation of sulfate assimilation loci of selected strains for construction of a phylogenetic tree (Fig. 3). Abbreviations: Sm, S. meliloti (NodP1, gi14523565; NodP2, gi15140612; CysD, gi5911360); Sbr, Sinorhizobium sp. strain BR816 (NodPl, gi2148989; NodP2, gi27125923; CysD, gi24528409); Rt, R. tropici CFN299 (NodP, gi1280528; CysD, gi7387610); Bm, Brucella melitensis (CysD, gi17988038); N33, Mesorhizobium sp. strain N33 (NodP, gi1531624); Ml, Mesorhizobium loti (NodP, gi13476292); Be, Bradyrhizobium elkanii (NodP, gi14209498); Ab, Azospirillum brasilense (NodP, gi142424); Ec, E. coli (CysD, gi12517206); Ka, Klebsiella aerogenes (CysD, gi11992146); Xo, X. oryzae pv. oryzae (NodP, gi21105248); Mt, Mycobacterium tuberculosis (CysD, gi15608425); Sc, Streptomyces coelicolor (CysD, gi21224427); At, Agrobacterium tumefaciens (CysD, gi15155798). S, fully sulfated Nod factors; S/NS, mixture of sulfated and nonsulfated Nod factors; NS, nonsulfated Nod factors; APR, APSreducing activity; PAPR, PAPS-reducing activity; ?APR, putative APR-reducing activity; ?, APS or PAPS reductase activity unknown; *, genome sequence not (fully) determined. Similar open reading frames are shaded identically. Note that nodP1 of S. meliloti is located on megaplasmid 1, nodP2 is on megaplasmid 2, and cysHDN is chromosomally located. nodP1 of Sinorhizobium sp. strain BR816 is located on a megaplasmid, nodP2 is on the symbiotic plasmid, and cysHDN is chromosomally located.

made for *S. meliloti* and *R. tropici*. In summary, these data support the ATP sulfurylase activity of the putative proteins encoded by the isolated BR816 cysDN genes.

Phylogenetic analysis of CysD and CysN homologues. The BR816 CysD and CysN ATP sulfurylase subunits were compared through multiple-sequence alignment (ClustalW) with homologous ATP sulfurylases subunits retrieved from Gen-Bank. The genomic organizations of the different sulfate assimilation loci of the strains selected for the phylogenetic analysis are schematically drawn in Fig. 2. Phylogenetic analysis of cysD and nodP gene products by the protein parsimony method resulted in a maximum-parsimony tree, as shown in Fig. 3. An identical tree topology could be inferred by using the neighborjoining method (data not shown). Similar phylogenetic relationships could be deduced after construction of a phylogenetic dendrogram of CysN and NodQ protein sequences by using either the neighbor-joining method or protein parsimony analysis (data not shown).

It can be observed that the CysD and NodP ATP sulfurylase subunits of *Rhizobium* spp. producing sulfated Nod factors

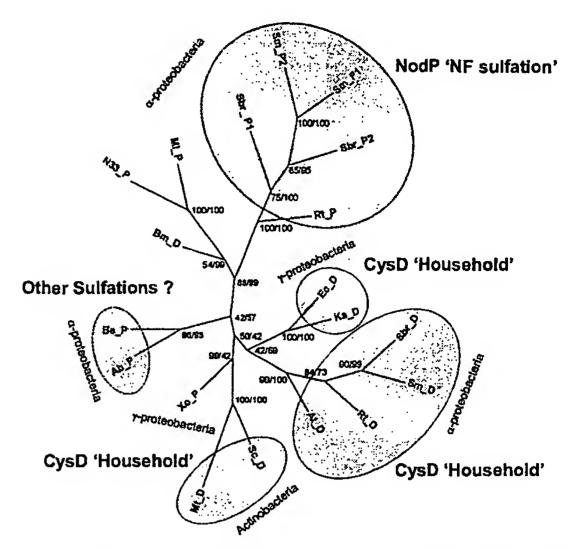


FIG. 3. Phylogenetic relationships among cysD gene products. The tree topology was inferred by using the protein parsimony method. Numbers represent the bootstrapping score (9) over 1,000 trials (parsimony/distance). The abbreviations of the species are as for Fig. 2.

(Fig. 2), which have been shown to be involved in amino acid biosynthesis (1, 23) and Nod factor sulfation (24, 25, 47), respectively, cluster in two different groups (Fig. 3). The CysD protein of *Sinorhizobium* sp. strain BR816 clearly belongs to the protein cluster involved in biosynthesis of sulfur-containing amino acids, supporting its putative function.

Two other "household" clusters could be distinguished, i.e., the γ-Proteobacteria clade and the Actinobacteria clade. Interestingly, only one gene copy coding for a sulfate activation complex has been described, for Mycobacterium tuberculosis (cysDNC) (Fig. 2) (58). The sulfate assimilation pathway of Mycobacterium tuberculosis proceeds from sulfate through APS (catalyzed by CysDN), which is converted by APS reductase (CysH) in the first step toward cysteine and methionine. APS can also be converted to PAPS, through the action of the APS kinase CysC, and serves as a substrate for sulfotransferases that produce sulfolipids, which putatively function as virulence factors (58). Similarly, APS and PAPS pools are generated through the enzymatic activity of RaxP and RaxQ in X. oryzae pv. oryzae and are used for both cysteine synthesis and sulfation of avirulence effector molecules (48).

The CysD-homologous proteins of some members of the Rhizobiaceae (among which are Mesorhizobium loti, producing nonsulfated Nod factors [29, 34]; Mesorhizobium sp. strain N33, producing sulfated Nod factors [35]; and the pathogen Brucella melitensis) seem to belong to another cluster. However, these proteins are still more closely related to the NodP Nod factor sulfation cluster than to the CysD household cluster, as defined above. Brucella melitensis was previously shown to be genetically closely related to Rhizobium spp. (14). Intriguingly, the respective Bradyrhizobium elkanii and Azospirilum brasilense ATP sulfurylase subunits constitute a separate cluster (Fig. 3). The nodPQ genes of B. elkanii are situated within a gene cluster comprising genes for symbiotic functions (fixGHIS and noeE) as well as genes involved in rhizobitoxin

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biosynthesis (59). Since the *B. elkanii* Nod factors are not sulfated (4, 43), these genes do not function in Nod factor biosynthesis. The recently finished genome sequencing of the p90 plasmid of *A. brasilense* sheds new light on a possible function of its *nodPQ* copy, which is located within a region carrying genes involved in polysaccharide synthesis (E. Vanbleu and J. Vanderleyden, unpublished results). It was previously shown that *A. brasilense* does not synthesize Nod factors and that deletion of the *nodPQ* copy does not lead to auxotrophy (54). Therefore, it can be speculated that this cluster encompasses proteins belonging to a novel functionality group.

Growth characteristics of Sinorhizobium sp. strain BR816 cysD mutants under free-living conditions. To investigate the biochemical role of the isolated cysDN genes of BR816, the BR816 cysD gene was mutated (see Materials and Methods). First, the cysD mutants were tested for cysteine auxotrophy. In addition, we were interested to know whether a cysD mutation could be complemented by one or both nodP copies of Sinorhizobium sp. strain BR816. Growth of the wild type and various mutants with mutations in nodPQ and/or cysDN (FAJ1600, FAJ1604, CFNE205, CFNE206, CFNE207, and CFNE208) was examined in liquid sulfate-free AMS medium supplemented with various sulfur sources (see Materials and Methods). It could be demonstrated that the BR816 nodPQ single or double mutants (CFNE205, CFNE206, CFNE207, and CFNE208) exhibit growth patterns similar to that of the wild-type strain in minimal medium with sulfate as the sole sulfur source (data not shown). Therefore, it can be concluded that nodPQ mutants are not auxotrophs. Growth of the cysD mutant (FAJ1600) with sulfate as the sole sulfur source was clearly affected compared to that of the wild-type strain (Fig. 4A). FAJ1600 showed a prolonged lag phase, although its generation time in exponential growth phase did not markedly differ from that of the wild type. The nodP1 nodQ2 cysD triple mutant (FAJ1604) was completely impaired in growth (Fig. 4A). In the presence of sulfite, cysteine, or methionine, the growth of both mutants after 60 h was nearly restored to the wild-type level (Fig. 4B to D). This indicates that the cysDN genes are effectively involved in the biosynthesis of sulfurcontaining amino acids, more specifically in the step of the sulfate assimilatory pathway just before the reduction of activated sulfate to sulfite. From this experiment we can conclude that knocking out the three sulfate activation systems (FAJ1604) in Sinorhizobium sp. strain BR816 leads to cysteine auxotrophy.

Interestingly, the growth characteristics of FAJ1600 showed a course similar to that of the wild type after a certain time interval. This demonstrates that the PAPS pool generated by the NodPQ sulfate activation complex is accessible for reduction by CysH and thus is available for the biosynthesis of sulfur-containing amino acids. The growth delay of FAJ1600 might indicate that CysH of *Sinorhizobium* sp. strain BR816 preferentially shows APS reductase activity rather than PAPS reductase activity toward the formation of sulfite. Moreover, the APS reductase activity of CysH has been recently confirmed in many rhizobial species (1, 20).

One should consider that (i) the growth curves of the wildtype and mutant strains were monitored under conditions in which no Nod factors are produced (no flavonoid induction) and (ii) nodP2, which is localized in the nodulation region on

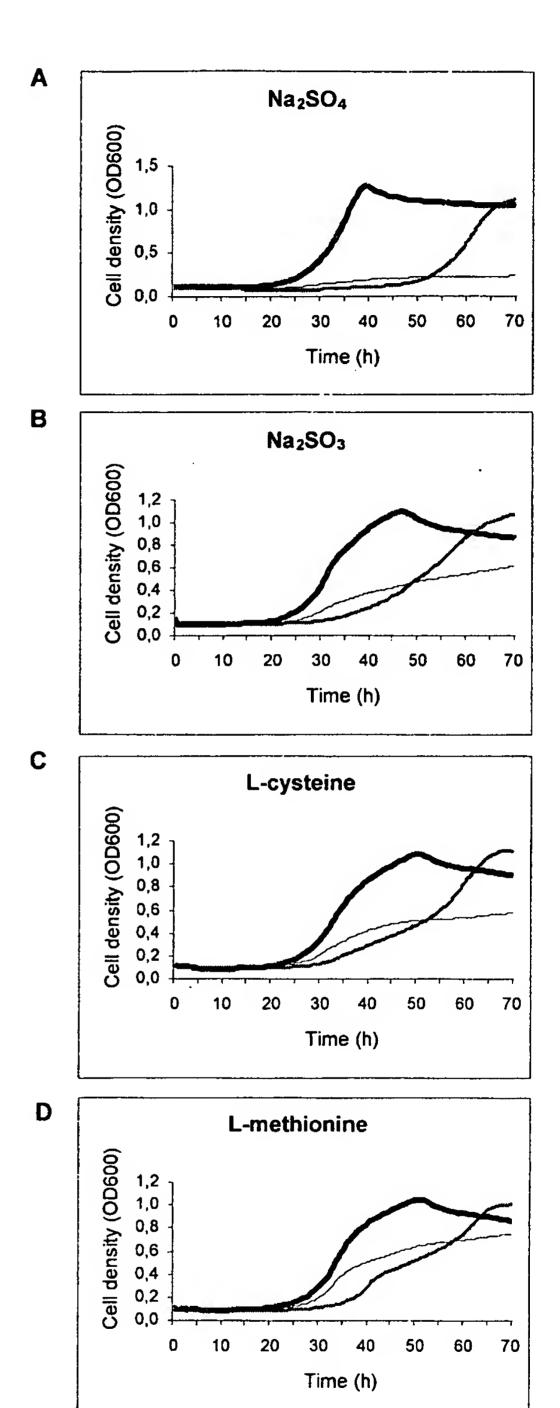


FIG. 4. Effects of various sulfur sources on cell growth of Sinorhizobium sp. strain BR816 wild-type and mutant strains determined by measuring optical density at 600 nm (OD600) in a BioscreenC instrument over a 4-day period. Thick black line, BR816; gray line, FAJ1600; thin gray line, FAJ1604). Cultures were grown at 30°C in sulfate-free AMS medium supplemented with sodium sulfate (A), sodium sulfite (B), L-cysteine (C), or L-methionine (D) at a concentration of 25 µM. Each experiment was conducted three times. Results from one experiment are shown.

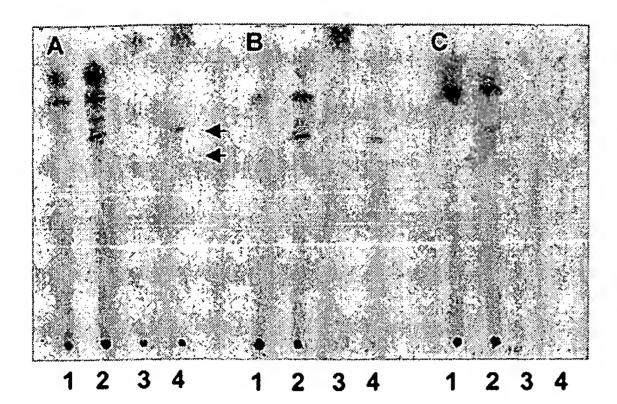


FIG. 5. Autoradiogram of a reverse-phase TLC profile of butanol extracts of radioactively labeled *Sinorhizobium* sp. strain BR816 (A), FAJ1600 (B), and FAJ1604 (C). Lanes 1 and 2, ¹⁴C labeling; lanes 3 and 4, ³⁵S labeling. Lanes 1 and 3, noninduced; lanes 2 and 4, apigenin induced. Spots representing sulfated Nod factors are indicated with arrows.

the symbiotic plasmid, probably is *nod* box dependent and thus not expressed (49). Therefore, to investigate whether the simultaneous production of sulfated Nod factors affects growth characteristics of the *cysDN* mutant strains, similar growth tests were performed in the presence of the *nod* gene inducer apigenin and with sulfate as the sole sulfur source. In this case, similar growth courses were obtained for FAJ1600 and FAJ1604 compared to the wild type (data not shown). This implies that at least the expressed *nodPQ* copy can complement and is sufficient for growth of FAJ1600 in minimal medium with sulfate as the sole sulfur source. The use of higher concentrations of inducer did not have a significant effect on the growth curves of the strains tested.

Nod factor sulfation pattern of Sinorhizobium sp. strain BR816 cysD mutants. Since the available nodPQ single and double mutants of Sinorhizobium sp. strain BR816 (CFNE205, CFNE206, CFNE207, and CFNE208 [Table 1]) were not auxotrophic and still produced sulfated Nod factors, Laeremans et al. (25) speculated that the housekeeping cysDN(C) genes can complement mutations in genes responsible for Nod factor sulfation. In order to determine to what level the Nod factors produced by the wild-type strain and the mutant strains FAJ1600 and FAJ1604 were still sulfated, apigenin-induced cell cultures, grown in liquid sulfate-free AMS medium supplemented with cysteine, were labeled with [14C]acetate or [35S]sulfate, and butanol extracts of the cell cultures were analyzed by reverse-phase TLC. Separation of the BR816 Nod factors revealed the presence of apigenin-induced spots on the chromatogram, corresponding to the Nod factors of BR816 (Fig. 5). The triple mutant FAJ1604 no longer produced sulfated Nod factors, which is in clear contrast with the sulfated Nod factor pattern of both the wild-type strain and FAJ1600 (Fig. 5). These results indicate that an activated sulfate source needed for Nod factor sulfation is no longer present. It can be concluded that the cysDN sulfate assimilation locus does provide active sulfate for NF sulfation.

Symbiotic phenotype of cysD mutants. The Sinorhizobium sp. strain BR816 cysD mutants were tested for their ability to

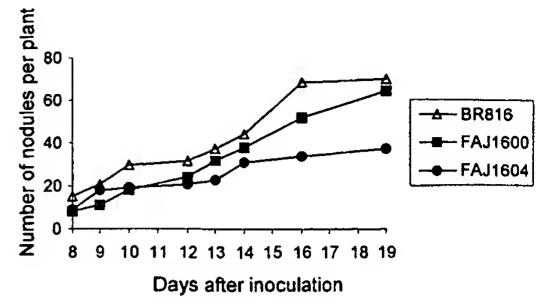


FIG. 6. Nodulation kinetics of *P. vulgaris* BAT477 inoculated with *Sinorhizobium* sp. strain BR816 wild-type and mutant strains. Two independent experiments were set up, and the results of one experiment are shown.

nodulate common bean (P. vulgaris ev. BAT477) and to fix nitrogen. No significant differences in the kinetics of appearance of the first nodules were observed (Fig. 6). However, FAJ1600 (cysD) as well as FAJ1604 (nodP1 nodQ2 cysD) showed a decreased nodule number per plant over time, but only for FAJ1604 was this difference significant at the 95% level (Tukey's test). Morphologically, the nodules of both mutant strains were generally smaller with apparently less leghemoglobin present (as judged by the absence of pink color).

To study the nitrogen fixation capacity of the nodulated roots, the acetylene reduction activity was measured. The acetylene reduction activity of 21-day-old nodules induced by FAJ1600 or FAJ1604 was significantly lower than that for the wild-type strain (P < 0.05; Tukey's test) (data not shown). When methionine was added to the plant nutrient solution, the nitrogen fixation per plant was restored to wild-type levels. Interestingly, supplementation with methionine resulted in an overall higher nitrogen fixation capacity of P. vulgaris cv. BAT477 inoculated with Sinorhizobium sp. strain BR816 (data not shown).

DISCUSSION

In this study, a third APS-producing locus of the broad-hostrange strain Sinorhizobium sp. strain BR816 was isolated. The nucleotide sequence of this region was determined, and based on homology searches, cysD and cysN were identified. Like in S. meliloti, no cysC homologue could be isolated downstream from cysDN. This is an indication that, like in other rhizobia, APS rather than PAPS is reduced to sulfite for cysteine biosynthesis (1). The highest similarity was found with the cysDN homologues in S. meliloti, supporting the close phylogenetic relationship between S. meliloti and Sinorhizobium sp. strain BR816 (15). Phylogenetic analysis revealed that CysD does not cluster with NodP1 and NodP2. The two BR816 NodP proteins are closely related and could have originated from a recent gene duplication, as was proposed for the NodP proteins of S. meliloti (13). Within the α -Proteobacteria clade, two clusters of proteins are clearly functionally distinguished and were designated NodP Nod factor sulfation and CysD household. It has been demonstrated that the nodPQ genes are also required for sulfation of S. meliloti lipopolysaccharide, proving a dual functionality of members of the NodP Nod factor sulfation cluster

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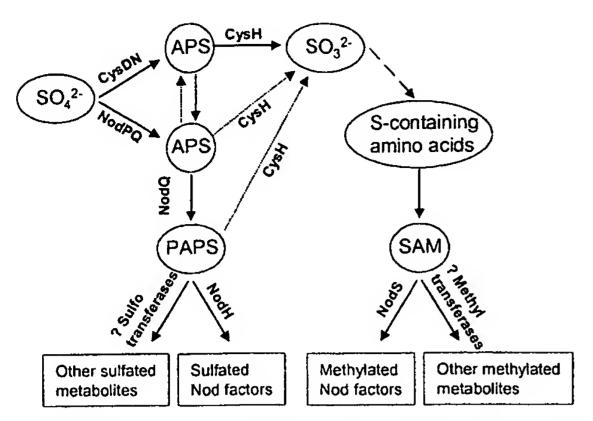


FIG. 7. Schematic representation of the distribution of APS and PAPS for sulfation and methylation processes in *Sinorhizobium* sp. strain BR816. Dotted arrows indicate possible but less favorable enzyme activity.

(5, 17). A potential new NodP-like protein cluster is proposed, comprising proteins involved in sulfate activation for sulfation of compounds that are yet unknown but which could be important during symbiosis. Other closely related CysD and NodP homologous do not fit into a specific functionality group, since these proteins are involved either in sulfation of amino acids (M. loti and B. melitensis) or in sulfation of Nod factors (Mesorhizobium sp. strain N33). It should be noted that within the γ-Proteobacteria clade and the Actinobacteria clade, only one copy of genes encoding sulfate activating enzymes is present, which seems to be involved in biosynthesis of sulfurcontaining amino acids as well as sulfation of other macromolecules.

We examined the effect of a cysD mutation under free-living conditions in a wild-type chromosomal background and in a nodP1 nodQ2 double mutant background. The levels of Nod factor sulfation (Fig. 5) as well as the growth characteristics (Fig. 4) of the different mutants were determined. In this study, we could demonstrate that the household CysDN sulfate activation locus of BR816 can additionally ensure Nod factor sulfation, whereas the symbiotic (P)APS pool, generated by the nodPQ sulfate activation complexes, can be engaged for sulfation of amino acids. Figure 7 shows a model of how the various forms of activated sulfate in Sinorhizobium sp. strain BR816 may be partitioned into the pathways for amino acid biosynthesis and sulfation of Nod factors and other compounds that might be important during symbiosis. The cysDN-dependent APS pool supplies activated sulfate that is subsequently reduced to form sulfite by the CysH APS reductase. Sulfite is further reduced to sulfide, which is then incorporated into the cysteine and methionine biosynthesis pathway. Our data suggest that the symbiotic APS and/or PAPS pool, created by the nodPQ-dependent sulfate activation step, can also be used by CysH (in a less efficient manner) for the biosynthesis of sulfurcontaining amino acids, when needed. Moreover, both household and symbiotic APS pools can be mutually exchanged. In S. meliloti, the nodPQ- and cysDN-encoded sulfate activation systems cannot substitute for each other (46, 47).

Why would Sinorhizobium sp. strain BR816 possess three

functional sulfate activation systems for Nod factor sulfation? Besides the use of activated sulfate for the biosynthesis of sulfur-containing amino acids and sulfation of Nod factors, (P)APS is needed for Nod factor methylation (37). Introduction of the S. meliloti nodPQ genes into R. tropici resulted in a decreased rate of R. tropici Nod factor methylation, while all R. tropici Nod factor backbones were sulfated. Waelkens et al. (57) showed that methylation of Nod factors is required for nodulation of bean. In Sinorhizobium sp. strain BR816, the three operational sulfate-activating systems could play an important role in maintaining substitutions of bacterial determinants for symbiosis.

An R. tropici nodPQ mutant (producing drastically reduced amounts of sulfated Nod factors) and an R. tropici nodH mutant (producing nonsulfated Nod factors) still activate the signaling cascade for emergence of effective nodules on P. vulgaris roots (12, 24). For bean plants, the sulfate moiety of the Nod factor was shown to be involved in the efficiency of nodule formation but appears not to be essential (11, 22). The effects of the cysD mutant FAJ1600 and the nodP1 nodQ2 cysD triple mutant FAJ1604 on bean symbiosis were seen mainly in the reduction of nodule number per plant. Since under free-living conditions, a cysDN-dependent biosynthesis of sulfur-containing amino acids is essential to allow optimal growth of Sinorhizobium sp. strain BR816 with sulfate as the sole sulfur source, bacterial growth defects are likely the main reason for the decreased nitrogen fixation of bean plants inoculated with the mutants FAJ1600 and FAJ1604. These defects can be restored by the addition of methionine to the plant nutrient solution. We propose that at the early stages of the nodulation, the plant root exudates of the germinated seedlings provide enough sources of organic sulfur to allow bacterial growth. However, a shortage of an organic sulfur source like methionine impairs bacterial growth inside the plant. Inoculation experiments with a Rhizobium etli metZ (O-succinylhomoserine sulfhydrylase for methionine biosynthesis) (51) mutant on bean plants resulted in the formation of ineffective (Nod+ Fix⁻) nodules, which suggested that root cells do not supply the inoculant bacteria with enough methionine. The fact that supplemented methionine resulted in an overall higher nitrogen fixation capacity of P. vulgaris BAT477 inoculated with BR816 strains supports this hypothesis. In contrast to our observations, an R. etli cysG (siroheme synthetase for cysteine biosynthesis) mutant, which is able to induce the formation of effective nodules (Nod+ Fix+) on the roots of common bean, seems to dispose of an organic sulfur source like cysteine or glutathione to allow growth inside the plant (52).

How can the strictly separated symbiotic and endogenous (P)APS pools in S. meliloti versus the complementary (P)APS pools in Sinorhizobium sp. strain BR816 be explained? Presumably, the nodPQ genes arose in ancestral rhizobial strains through duplications of the endogenous cysDNC genes. Later, these nodPQ genes evolved toward more specialized symbiotic genes, whereas the endogenous cysC gene, encoding the APS kinase, was apparently lost during evolution. At this stage, complementation between both PAPS pools was still possible (the case of Sinorhizobium sp. strain BR816). Then, the genetic separation of the two sulfate-activating systems could have further evolved into two more efficient and energy-saving sep-

arate enzymatic multienzyme complexes (the case of S. meliloti).

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General information about the entry				
Entry name	IMP_1			
Accession number	PS00629			
Entry type	PATTERN			
Date	JUN-1992 (CREATED); APR-2006 (DATA UPDATE); JUL-2006 (INFO UP			
PROSITE documentation	PDOC00547			
The second secon	Name and characterization of the entry			
Description	Inositol monophosphatase family signature 1.			
Pattern	[FWV]-x(0,1)-[LIVM]-D-P-[LIVM]-D-[SG]-[ST]-x(2)-[FYA]-x(0,1)-[HKRNSTY			
	Numerical results			

- UniProtKB/Swiss-Prot release number: 50.4, total number of sequence entries in that re 230133.
- Total number of hits in UniProtKB/Swiss-Prot: 75 hits in 75 different sequences
- Number of hits on proteins that are known to belong to the set under consideration: 74
 different sequences
- Number of hits on proteins that could potentially belong to the set under consideration:
 different sequences
- Number of false hits (on unrelated proteins): 0 hits in 0 different sequences
- Number of known missed hits: 5
- Number of partial sequences which belong to the set under consideration, but which are pattern or profile because they are partial (fragment) sequences: 1
- Precision (true hits / (true hits + false positives)): 100.00 %
- Recall (true hits / (true hits + false negatives)): 93.67 %

Comments

- Taxonomic range: Archaebacteria, Eukaryotes, Prokaryotes (Bacteria)
- Maximum known number of repetitions of the pattern in a single protein: 1
- 'Interesting' site in the pattern: 4, metal(?)
- Interesting' site in the pattern: 9,metal(?)
- VERSION: 2

Cross-references

True positive hits:

BPNT1 HUMAN	(095861),	BPNT1_MOUSE	(Q9Z0S1),	BPNT1_RAT
CYSQ ACTAC	(P70714),	CYSQ_BUCAI	(P57624),	CYSQ_ECO57
CYSQ ECOL6	(Q8FAG5),	CYSQ_ECOLI	(P22255),	CYSQ_HAEIN
CYSQ MYCBO	(P65164),	CYSQ_MYCLE	(P46726),	CYSQ_MYCTU

```
CYSQ SALTI (Q8Z153), CYSQ_SALTY (P26264), CYSQ_SHIFL
              DPNP1_ARATH (Q42546), DPNP2_ARATH (O49623), DPNP4_ARATE
              DPNPH ARATH (Q38945), DPNPM ARATH (Q9M0Y6), DPNP_ORYSA
                           (094505), HAL21_CANAL (P46594), HAL22 CANAI
              DPNP SCHPO
              HAL2 YEAST (P32179), IMP1 LYCES
                                                   (P54926), IMP2 LYCES
              IMP3 LYCES (P54928), IMPA1 BOVIN (P20456), IMPA1 CAEEI
              IMPA1 HUMAN (P29218), IMPA1 MOUSE (055023), IMPA1 PIG
              IMPA1 PONPY (Q5R4X0), IMPA1 RAT
                                                   (P97697), IMPA1 XENLA
              IMPA2 HUMAN (014732), IMPA2 MOUSE (Q91UZ5), IMPA2 RAT
              IMPP MESCR (049071), INM1 YEAST
                                                   (P38710), INM2 YEAST
              INPP BOVIN (P21327), INPP HUMAN
                                                   (P49441), INPP MOUSE
              QAX NEUCR
                           (P11634), QUTG EMENI
                                                   (P25416), SUHB AQUAE
              SUHB ARCFU
                           (030298), SUHB_BACSU
                                                   (Q45499), SUHB_CAUCR
                           (POADG6), SUHB ECOL6
                                                   (POADG5), SUHB ECOLI
              SUHB ECO57
                          (P44333), SUHB METJA
                                                   (Q57573), SUHB METTH
              SUHB HAEIN
                           (P65166), SUHB MYCLE
                                                   (P46813), SUHB MYCTU
              SUHB MYCBO
                          (Q9JU03), SUHB NEIMB
                                                   (Q9JZ07), SUHB PASMU
              SUHB NEIMA
                           (Q9HXI4), SUHB RHILO
                                                   (Q98F59), SUHB RHIME
              SUHB PSEAE
                           (P58537), SUHB SYNY3
              SUHB SALTY
                                                   (P74158), SUHB THEMA
                           (Q9KTY5), SUHB XYLFA
                                                   (Q9PAM0), SUHB XYLFT
              SUHB VIBCH
UniProtKB/Swiss-Y4FL_RHISN
                           (P55450), YHEB_CHLVI
                                                   (P56160)
Prot
              False negative hits (sequences which belong to the set under conside
              which have not been picked up by the pattern or profile):
              DPNP3 ARATH (Q8GY63), SUHB AERPE
                                                   (Q9YAZ7), SUHB_BUCAI
              SUHB BUCAP (Q8K9P6), SUHB BUCBP
                                                   (Q89AK9)
              `Potential' hits (partial sequences which belong to the set under cons
              which are not hit by the pattern or profile because they are partial (fra
              sequences):
              YPSS RHILP
                          (P10497)
              Sequences which could potentially belong to the set under considera
              PPNK METJA
                           (Q58327)
              Retrieve an alignment of UniProtKB/Swiss-Prot true positive hits:
              [Clustal format, color, condensed view] [Clustal format, color] [Clustal form
              [Fasta format]
              1AWB; 1DK4; 1FLF; 1G0H; 1G0I; 1IMA; 1IMB; 1IMC; 1IMD; 1IME; 1IMF;
PDB
              1K9Y; 1K9Z; 1KA0; 1KA1; 1LBV; 1LBW; 1LBX; 1LBY; 1LBZ; 1Q00; 1QGX
[Detailed view]
              2HHM;
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